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(54) **Intermediates in the synthesis of vitamin D derivatives.**

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**JOURNAL OF ORGANIC CHEMISTRY, vol. 41,
no. 12, 11th June 1976, pp. 2098-2102,
Washington D.C. (US); D. JOHN ABERHART et
al.: "Studies on the adduct of 4-phenyl-1,2,4-
triazoline-3,5-dione with Vitamin D3"**

**CHEMICAL ABSTRACTS, vol. 92, no. 19, 12th
May 1980, p. 624, no. 164161d, Columbus, Ohio
(US); E.ZBIRAI et al.: "Structural
transformations on vitamin D"**

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Description

This invention relates to novel intermediates in the production of vitamin D analogues and of vitamin D analogues which may be produced therefrom.

In the past modified vitamin D derivatives have been prepared through modification of sterol precursors which are then converted into vitamin D derivatives through a standard series of steps, normally preliminary conversions of $\Delta^{5,7}$ dienes followed by irradiation of the dienes to give D vitamins. These procedures have serious flaws. First, all of the available methods for the synthesis of $\Delta^{5,7}$ dienes tend to give mixtures of products or require a number of steps and proceed in relatively low yield. The second difficulty is that the only known transformation of the 5,7 dienes into the vitamins involves irradiation followed by thermal equilibration. Irradiation intrinsically gives rise to a mixture of byproducts. This limits the yield of the desired vitamin and furthermore complicates its recovery in pure form.

Previous attempts to modify the 17-side chain of vitamin D compounds have been unsuccessful due to instability problems. We have now found that vitamin D₂ and related compounds can be converted to a protected form capable of withstanding the reaction conditions necessary for oxidative cleavage of the 22,23-double bond to form a 22-aldehyde which can then be converted to other derivatives as described hereinafter. In particular, we have found that vitamin D₂ compounds in either the *cis* or *trans* configuration can be stabilised by formation of a Diels Alder dienophile adduct which can subsequently be reconverted to the *trans* form of the vitamin after the side-chain modification. The *trans* vitamin analogues can then be efficiently converted into the active *cis* form by known reactions.

The formation of certain dienophile adducts of vitamin D₃ has been described in the literature. D. J. Aberhart and A. Chi Tung Hsu (J. Org. Chem. Vol. 41, No. 12, 1976, 2098-2102) have described the formation of an adduct with 4-phenyl-1,2,4-triazoline-3,5-dione and E. Zbiral and W. Reischl (Proc. Workshop Vitamin D 1979, 4th. (Vitam. D. Basic Res. Its Clin Appl.), 21-24) have also described the adduct with sulphur dioxide. However, there is no disclosure of the use of these dienophiles to form adducts with vitamin D₂ followed by oxidative cleavage at the 22,23-double bond while leaving double bonds in the 5,10- and 7,8-positions intact.

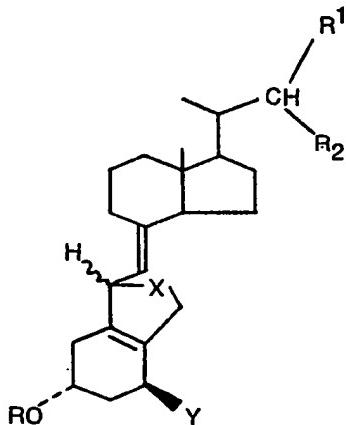
According to one feature of the present invention we provide compounds of the general formula I,

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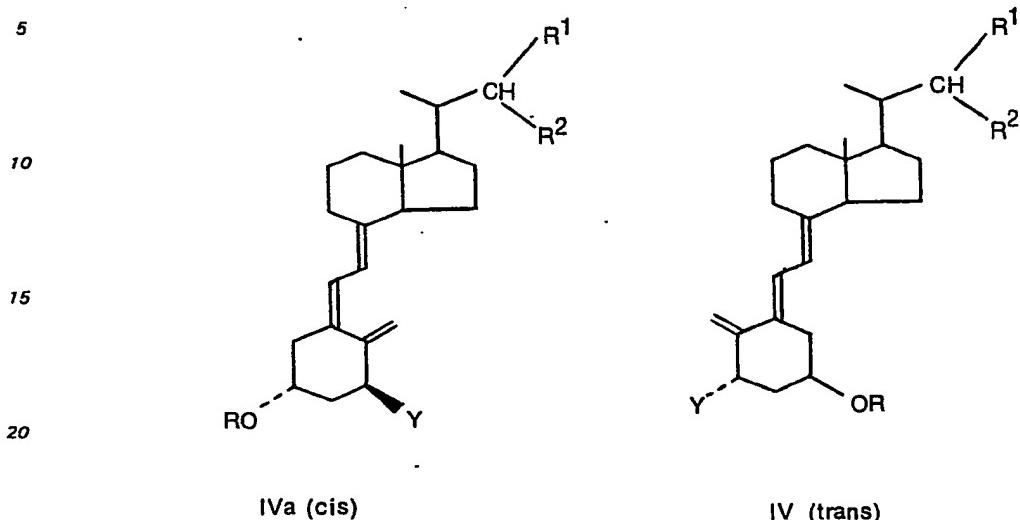


wherein R represents a hydrogen atom or a hydroxyl protecting group, Y represents a hydrogen atom or an optionally protected hydroxyl group, X represents $-\text{SO}_2-$ or the residue of a diacylazo dienophile and either R¹ represents a halogen atom a hydrocarbysulphonyloxy group or a group of the formula $-\text{Z}-\text{R}^3$ (in which Z represents $-\text{O}-$, $-\text{S}-$, $-\text{SO}-$, $-\text{NR}^4-$ or $-\text{CR}^4\text{R}^5-$ and R³, R⁴ and R⁵, which may be the same or different, each represents a hydrogen atom or a straight or branched aliphatic group having 1-12 carbon atoms and which may optionally carry one or more substituents) and R² represents a hydrogen atom or R¹ and R² together represent an oxo group or an optionally substituted alkylidene group, except that R¹ together with the group $-\text{CH}(\text{CH}_3)\text{CH}-$ to which they are attached do not represent a group having the branched 17beta-hydrocarbyl side chain skeleton of vitamin D₂ or vitamin D₃.

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The above compounds are useful intermediates in the preparation of vitamin D analogues i.e. compounds of general formula IV and IVa.



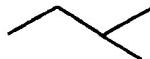
wherein R, Y, R¹ and R² are as hereinbefore defined. The above compounds of general formulae IV and IVa are also novel and constitute a still further feature of this invention.

The use of the compounds of general formula I in the preparation of the novel compounds of formulae IV and IVa is illustrated in the reaction scheme of the accompanying drawings, R, Y, X, R¹ and R² being as defined above. The compounds of formula I—IV may also carry further groupings.

It should be noted that the Diels Alder adduct formed from either the 5,6-*cis*- or the 5,6-*trans*-vitamin starting material exists as a mixture of two possible isomers at the 6-position. However, since the eventual removal of the Diels Alder residue always generates a compound of the 5,6-*trans* configuration, there is no need to distinguish between such 6-isomers or to effect their separation.

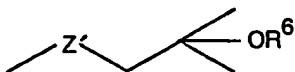
We have found that using the above procedure a wide range of groups R¹ may be introduced into the vitamin D structure. Thus, as indicated above R¹ may be a group of the formula Z—R³, where Z is —O—, —S—, —SO—, —NR⁴ or —CR⁴R⁵— and R³, R⁴ and R⁵, which may be the same or different, are each a hydrogen atom or a straight or branched aliphatic group having 1—12 carbon atoms which may carry one or more substituents such as, for example halogen atoms (e.g. fluorine), or optionally protected hydroxyl groups.

In general it is preferred that the group R^3 in the final products should be of the formula



(in order to provide a 17 β -side chain of approximately the shape present in natural vitamin D compounds) with the possibility of substitution as described above. The heteroatoms Z, where present, do not greatly change the overall shape of the side chain.

In particular, the invention enables compounds of formula IV and IVa to be prepared in which R¹ is of formula



wherein Z' represents —O—, —S—, —NH— or —SO— and R⁶ represents a hydrogen atom or a hydroxyl protecting group, the 1a-position optionally carrying a hydroxyl or protected hydroxyl group, these being analogues of the active metabolite 25-hydroxyvitamin D³.

Protected hydroxyl groups may, for example, be acyl groups e.g. alkanoyl groups (preferably having 1—6 carbon atoms), aralkanoyl groups (preferably having 7—15 carb n atoms), aroyl groups (preferably having 6—12 carbon atoms), cyclic ether groups or tri-hydrocarbysilyl groups. Examples of such gr ups include acetyl, propionyl, benzoyl and tetrahydropyranyl groups and trihydrocarbysilyl groups having up to three C₁₋₆ alkyl, C₆₋₁₂ aryl and/or C₇₋₁₅ aralkyl groups.

The new synthetic analogues of the invention have modified vitamin D properties of interest in medicine.

The compounds of formulae IV and IVa in which R¹ has the above meanings may be prepared, *inter alia*, by nucleophilic substitution reactions on compounds of formula IV and IVa in which R¹ represents a halogen atom, such as a chlorine, bromine or iodine atom, or a leaving group, for example a hydrocarbonylsulphonyloxy group O—SO₂R⁷ in which R⁷ represents, for example, an alkyl group (preferably having 1–6 carbon atoms), an aryl group (preferably having 7–15 carbon atoms). The tosyloxy group is preferred. Alternatively, the above compounds may be prepared from corresponding compounds of formula I and the dienophile group X removed subsequently. Since, however, the nucleophilic substitution reactions are mostly carried out in the presence of a base and since the protected compounds of formula I are less stable to base than the parent trienes of formula IV and IVa, the latter are commonly preferred substrates.

In the formation of 22-thia compounds (in which Z is —S—), the nucleophilic reagent is conveniently the thiol of formula R³XX reacted in an inert solvent such as tetrahydrofuran in the presence of a non-nucleophilic base, for example an inorganic base such as sodium hydride or an organic base such as pyridine.

The corresponding sulphoxides (Z=—SO—) may be prepared by oxidation of the thia-compound (Z=—S—), for example using a peracid or salt as oxidising agent, e.g. a periodate. Mixtures of the (R) and (S) sulphoxides are normally formed and the invention extends to these separately and in admixture.

The 22-oxa compounds of formula IV or IVa may be prepared by reaction of an alcohol of formula I, IV or IVa in which R¹ is OH, with an alkylating agent or alternatively by reaction of a reactive derivative thereof, with an alcoholate; these reactions are followed by deprotection when a compound of formula I is used. The reactive derivative may, for example, be a halide such as an iodide, or a hydrocarbonylsulphonyloxy derivative, such as a tosyloxy derivative, the alcoholate being, for example, an alkali metal or thallium alcoholate of the alcohol R³OH. It is preferred, however, to react the compound of formula I, IV or IVa in which R¹ is OH with an epoxide. This generates a side chain carrying a hydroxyl group derived from the epoxide oxygen. Where it is desired to make 25-hydroxy-22-oxa vitamin D₃ derivatives, a suitable reagent is isobutylene epoxide.

The reaction is advantageously effected in an inert solvent, e.g. a hydrocarbon solvent such as benzene, in the presence of a non-nucleophilic base, conveniently an alkali metal t-alkoxide in the presence of a phase transfer agent such as a crown ether. Under such basic conditions, we have found it especially preferred to effect the reaction on a starting compound of formula IV or IVa, since the trienes are, as indicated above, more stable to these conditions than the dienophile-protected compounds of formula I.

The 22-aza compounds of formula I, IV or IVa may be prepared by reaction of a reactive derivative of an alcohol of formula I, IV or IVa in which R¹ is OH, for example a halide such as an iodide, or a hydrocarbonylsulphonyloxy derivative such as a tosyloxy derivative, with an amine of formula R³R⁴NH. Due to the basicity of the reagent, a substrate of formula IV or IVa is preferred. Where the amine is liquid it is preferably reacted without added solvent.

The 22-aza derivatives may often conveniently be isolated as N-acylates, such as N-acetates, which may be formed by reaction with an appropriate acid anhydride.

The 22-hydrocarbonylsulphonyloxy derivatives of formulae I, IV and IVa used in the above reactions may be prepared by reacting the corresponding alcohol with the appropriate hydrocarbonylsulphonyl halide, e.g. tosyl chloride in the presence of a base such as pyridine. Best results have been obtained by effecting this reaction on a compound of formula I in which X is SO₂, and removing the SO₂ residue by thermolysis, as described hereinafter.

The compounds of formula I, IV or IVa in which Z in R¹ is CR⁴R⁵ may be prepared by reacting compounds of formula I, IV or IVa carrying a hydrocarbonylsulphonyloxy group R¹, e.g. a tosyl group, with carbon nucleophiles. Suitable carbon nucleophiles are Grignard reagents reacted in the presence of a copper catalyst, e.g. a cuprous salt. Thus, for example, 25-hydroxy vitamin D₃ and the 1α-hydroxy derivative thereof may be prepared by reacting an appropriate hydrocarbonylsulphonyloxy derivative of formula I, IV or IVa with a Grignard reagent of the formula



(where R⁶ has the above meaning) in tetrahydrofuran in the presence of cuprous iodide.

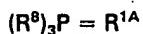
For the production of an alcohol of formula I in which R¹ is OH, for use in the preparation of the above novel vitamin D derivatives, the formyl group in the corresponding aldehyde of formula I (wherein R¹ and R² together represent oxo) must be reduced.

We have found that this can be effected readily, often in essentially quantitative yield, by reaction with a metal hydride reducing agent such as an alkali metal borohydride, e.g. sodium borohydride. It is noteworthy that this reduction retains the original configuration at the 20-carbon atom. Such alcohols are also new compounds.

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Compounds of formula I, IV or IVa may also be prepared in which R¹ is a divalent alkylidene group, which may carry substituents as described above for R³. Thus, for example, the aldehyde of formula I (wherein R¹ and R² together represent oxo) may be reacted with an ylid, for example a Wittig reagent which may be represented by the general formula

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wherein the groups R⁸, which may be the same or different, are alkyl (preferably C₁₋₆), aralkyl (preferably C₇₋₁₅) or aryl (preferably C₆₋₁₂) groups and R^{1A} is an alkylidene group (preferably having 1 to 8 carbon atoms) and may carry substituents as described for R³ above.

10 The Wittig reagent will normally be formed *in situ* by reaction of a quaternary salt thereof with a strong base in an inert solvent. Suitable bases include hydrocarbyl lithium compounds such as phenyl lithium and *n*-butyl lithium. Suitable solvents include ether solvents such as tetrahydropyran and diethyl ether. The aldehyde of formula I is preferably added immediately after the Wittig reagent has been formed.

15 The phosphonium salt precursor of the appropriate Wittig reagent for formation of the correct 17β-side chain of 25-hydroxy vitamin D₃ may, for example be prepared by reaction of isobutylene epoxide with methylenetriphenylphosphorane; the initially formed product in which R⁵ is H may if desired be protected, for example by formation of a tetrahydropyranyl or trihydrocarbysilyl derivative. The phosphorane is 20 preferably prepared by reaction of methyltriphenylphosphonium bromide in a cyclic ether solvent such as tetrahydrofuran in the presence of a strong base such as phenyl or *n*-butyl lithium, the isobutylene epoxide then being reacted *in situ* with a second equivalent of base. We have found the phosphonium bromide initially produced to be difficult to isolate and purify but that conversion to a tetraphenylborate salt enabled a relatively pure product to be obtained.

25 If it is desired to form a saturated side-chain, selective reduction of the newly formed 22,23-double bond is required. This was unexpectedly found to be possible using hydrogenation over 5% palladium on charcoal. It is noteworthy that this reduction restores the desired, "natural" configuration at the 22- and 23-carbon atoms. This route thus provides a further method of preparing compounds of formula I, IV or IVa in which R¹ is —CR³R⁴R⁵, as defined above.

30 It will be seen that the compounds of formula I in the above reaction scheme are key intermediates in the production of the new vitamin D analogues according to the invention. By way of illustration their preparation is now described in detail starting from vitamin D₂ or its 5,6-trans isomer.

The compound of formula III may be prepared by reaction of a vitamin D₂ compound of formula IIa or IIb with SO₂ or a diacylazo dienophile whereby the desired divalent grouping X is introduced.

35 Preferred diacylazo dienophiles are cyclic azo compounds such as phthalazine diones or triazoline diones; in general these may be represented by the formula V,

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where W is a divalent aromatic carbocyclic group such as a 1,2-phenylene group or a group

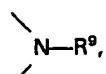
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50 where R⁹ is an aryl group such as a phenyl group. The divalent aromatic group or the aryl group R⁹ may carry substituents, for example C₁₋₆ alkyl or alkoxy groups, halogen atoms or nitro groups. Derivatives of formula III in which X is of formula V are also new compounds.

55 Where the dienophile is SO₂, this may simply be reacted with the vitamin D₂ compound in a suitable solvent, for example aqueous media capable of dissolving the vitamin. A well stirred mixture of water and a hydrocarbon solvent such as benzene is particularly useful. Basic conditions are preferably used, e.g. using an inorganic base such as an alkali metal bicarbonate. Where the dienophile is a cyclic azo compound of formula V in which W is a group

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65 this may be added to the starting vitamin D₂ compound in solution in a suitable solvent such as thyl acetat. Where W is a divalent-1,2-arylene grouping as in phthalazine 1,2-dione, however, this is preferably formed *in situ* by oxidation of the corresponding cyclic hydrazide, e.g. phthalhydrazide. Thus the vitamin D₂

compound may be reacted in solution in an inert solvent such as a halogenated hydrocarbon with the cyclic hydrazide in the presence of an oxidising agent such as lead tetraacetate/acetic acid.

After formation of the adduct of formula III, the 22,23-double bond may be cleaved to form the 22-formyl derivative of formula I by known oxidative techniques such as permanganate/periodate, osmate/periodate or, most preferably ozonolysis. We have found that this reaction proceeds selectively in high yield with little cleavage of the 7,8-double bond and, in particular, with no disturbance of the stereochemistry at the 20-position.

Ozonolysis may be effected by passing ozone, preferably diluted with a further gas such as oxygen, through a solution of the compound of formula III in a solvent therefor to form an ozonide which is then reductively cleaved by a suitable reducing agent. A suitable solvent is, for example, a halogenated hydrocarbon such as dichloromethane, a ketone, e.g. methyl ethyl ketone or acetone or an alcohol such as methanol or ethanol. A mixture of dichloromethane and methanol gave especially good yields. The reducing agent may be present during the reaction or added after ozonide formation is completed. Thus, for example tetracyanoethylene may be present in solution in acetone during ozonolysis. While reducing agents such as dimethyl sulphide may be used to reduce the ozonide after its formation, preferred reagents are trivalent phosphorus compounds such as triphenylphosphine.

Where an alcohol solvent is used, the aldehyde product of formula I may form an acetal derivative with the alcohol. This may, however, readily be cleaved hydrolytically, for example using an aqueous base e.g. sodium bicarbonate. The reaction is preferably carried out at low temperatures, for example, -78°C.

After modification of the 17-side chain, the residue X may be removed to yield, as indicated above, a 5,6-trans vitamin of general formula IV. The removal of the residue X will be effected in different ways, depending on its nature.

Where X is SO₂, it is conveniently removed by thermolysis under basic conditions, e.g. in the presence of a hydroxylic solvent such as an alcohol, e.g. ethanol, containing a base such as an alkali metal carbonate, e.g. sodium carbonate.

Where X is a group of formula V, removal can readily be effected by removal of the —CO—W—CO— moiety, for example by basic hydrolysis or treatment with hydrazine, followed by mild oxidation of the unsubstituted vitamin hydrazide so formed to the corresponding azo-compound which spontaneously decomposes to yield the required 5,6-trans vitamin. Basic hydrolysis can be effected using strong alkali such as sodium or potassium hydroxide, for example in solution in an alcohol such as methanol, or by treatment with an amine such as triethylamine. The preferred method, however, is treatment with hydrazine which produces the desired hydrazide in high yield; this reaction has not previously been described for decomposition of such Diels-Alder adducts. Oxidation may be effected using reagents capable of oxidising hydrazone compounds to azo compounds, for example ceric, cupric, ferric, ferricyanide or periodate salts or air. A preferred mild reagent, however, is a diaryl telluroxide such as dianisyl telluroxide, preferably used with a reoxidant such as 1,2-dibromotetrachloroethane and a base such as K₂CO₃, as described in our British Patent Specification No. 2058758A.

Where a 1α-hydroxy vitamin D compound is required, the modified 5,6-trans-vitamin compound of formula IV, which carries the desired 17-side chain, optionally protected, may be subjected to 1α-hydroxylation, using the procedure of our South African patent No. 79/5958. Thus, the 5,6-trans vitamin compound may be reacted with a selenite ester, preferably formed *in situ* by reaction of selenium dioxide and an alcohol such as methanol. The quantity of selenium compound may be reduced if a re-oxidant is employed, for example a periodate salt or N-methyl morpholine 1-oxide.

Alternatively, a reactive derivative of a 22-hydroxy derivative of formula I or IV above may be 1α-hydroxylated by the above procedure and the desired side-chain built up subsequently.

The 5,6-trans vitamin D compound of formula IV, after modifications such as those described above, may readily be isomerised in high yield to a required active cis-vitamin compound by known techniques, for example by irradiation in the presence of iodine or diphenyl selenide or, preferably, a triplet photosensitizer having a triplet energy of the order of 45±5 Kcal per mole, such as anthracene, acridine or phenazine. To avoid isomerization to undesired tachysterol derivatives, acid conditions should be avoided and the photoisomerisation is preferably effected in the presence of a base such as triethylamine.

Where protected hydroxyl groups are present in the vitamin product, these may be removed by conventional methods. In general, the vitamin structure is somewhat sensitive to acids, but is resistant to basic conditions and the latter are advantageously used. Acyloxy groups can thus be removed using alkali metal hydroxide in an alcohol solvent such as methanol. Silyl groups may be removed by treatment with quaternary ammonium fluorides such as tetra-n-butylammonium fluoride. Since most of the reactions described above can be applied to compounds having unprotected hydroxyl groups, protecting groups may be removed, if desired, at various stages. Although the vitamins are resistant to bases (and sensitive to acids), the dienophile adducts tend to be sensitive to bases and relatively resistant to acids. Consequently, acid conditions may be used to deprotect hydroxyl groups at stages where the dienophile residue X is present.

In general, most of the stages described above proceed in excellent yield. When conditions are optimised, yields of the order of 80% or more at each stage have been achieved. This renders the overall yield of modified vitamin, starting from vitamin D₂, markedly better than those achieved using many previously suggested routes.

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The following Examples are given by way of illustration only:—

Microanalyses and mass spectra were obtained by the staff at the Institut de Chemie des Substances Naturelles du CNRS, Gif-sur-Yvette, France. Melting points were determined using either a Kofler block, M I-temp or Fisher-Johns apparatus and are uncorrected. Optical rotations were measured at room temperature using a Rudolph Photoelectric Polarimeter, Model 70, and are reported for chl refrom solutions unless otherwise stated. UV spectra were recorded using a Carey 11 spectrophotometer and are reported for ethanol solutions. The molar extinction coefficient (ϵ) for these absorbances are given in parenthesis. IR spectra were recorded using a Perkin-Elmer 137 "Infracord" spectrophotometer and are reported for KBr discs unless otherwise stated. Absorbance characteristics are denoted by s = strong, m = medium, w = weak, sh = shoulder, br = broad. $^1\text{Hnmr}$ spectra were determined at 60MHz on a Varian T-60 spectrometer. NMR characteristics are denoted as s = singlet, d = doublet, tr = triplet, q = quartet, m = multiplet, W = peak width at half height and are reported for CDCl_3 solutions, unless otherwise indicated, with tetramethylsilane as internal standard, as values of δ (ppm downfield of TMS).

Thin layer chromatography (tlc) was carried out on 250μ silica gel GHLF "Uniplates" (Analtech, USA); and preparative layer chromatography (plc) on 1 mm silica gel GF-254 "Uniplates" (Analtech, USA). "Chromatography" refers to medium pressure liquid chromatography carried out using E. Merck silica gel 60H. High performance liquid chromatography (HPLC) was carried out using Waters Associates silica gel "Porasil A" packed in two 2 ft \times 3/8 inch stainless steel columns, and a Waters Associates chromatograph, equipped with a 6000 psi pump and a differential refractometer detector. Ozone was generated from a Towers Ozone Apparatus GE-150. Selective ozonolysis requires vigorous mixing of the dissolved substrate and the oxygen-ozone gaseous mixture. A "Vibromixer" (Chemapag, Switzerland) equipped with a stainless steel gas inlet/stirrer was particularly useful for this purpose. This equipment was also used for the formation of the phthalazine-1,4-dione Diels-Alder adducts of vitamin D.

A 200W Hanovia medium pressure mercury vapour lamp (654A36) was used as irradiation source for 5,6-double bond photoisomerisation reactions.

Reactions on calciferol substrates were routinely performed under an inert, argon atmosphere. Calciferols were stored at -20°C , under argon, in the dark, as either crystalline solids or (where possible) ether solutions. Solvents used were reagent grade unless otherwise stated.

Aqueous work-up refers to partition between an organic solvent and water, followed by sequential washing with a 5% aqueous sodium bicarbonate solution and a saturated aqueous sodium chloride solution. The organic solution was dried using either anhydrous MgSO_4 or anhydrous Na_2SO_4 , and the solvent removed on a rotary evaporator. Acid work-up refers to partition between an organic solvent and water, followed by sequential washing with a 4% aqueous HCl solution; 5% aqueous sodium bicarbonate solution, etc. as for aqueous work-up.

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Example 1

(a) 6(R),19-[4'-phenyl-1',2',4'-triazolidine-3',5'-dione-1',2'-yl]-9,10-seco- 3β -hydroxy-ergosta-5(10),7(E), 22(E)-triene
To ergocalciferol (5 g) in ethylacetate (150 ml) at 0°C under an argon atmosphere, 4-phenyl-1,2,4-triazoline-3,5-dione (2.4 g, 1.1 eq) in ethyl acetate (150 ml) was added over 45 min. After a further 1 hr, some of the title adduct had precipitated. The mixture was filtered and the filtrate passed down a neutral alumina column. Elution with hexane/ethylacetate gave the remainder of the product. Crystallisation from alcohol gave 6.2 g (86%), mp 99°C; $[\alpha]_D = +20^\circ$ ($c = 0.76$); $^1\text{Hnmr}$ δ 7.48 (s, 5H, aryl), 5.22 (m, W=10Hz, C—22H, 23H), 4.98 and 4.73 (an AB system, J=10Hz, C—6H, 7H), 4.2 and 3.85 (an AB system, J=15Hz, C—19H₂), 4.1 (m, C—3H), 0.533 (s, C—18H₃). IR v_{max} (CHCl_3) 3700 (br), 2950 (s), 1775 (m), 1710 (s), 1425 (s)cm⁻¹; mass spec. molecular ion, m/e = 571; (analysis found: % C, 75.63; H, 8.62; N, 7.36; $\text{C}_{38}\text{H}_{49}\text{O}_3\text{N}_2$; requires: % C, 75.62; H, 8.64; N, 7.35).

Similarly prepared from ergocalciferol acetate in 85% yield was the corresponding acetate 6(R),19-[4'-phenyl-1',2',4'-triazolidine-3',5'-dione-1',2'-yl]-9,10-seco- 3β -acetoxy-ergosta-5(10),7(E), 22(E)-triene.

Crystallised from ethanol.m.p. 85°C; $[\alpha]_D = +183^\circ$ ($c = 0.82$); $^1\text{Hnmr}$ δ 7.48 (s, 5H, aryl), 5.22 (m, W=12Hz, C—3H, 22H, 23H), 4.98 and 4.73 (an AB system, J=10Hz, C—6H, 7H), 4.2 and 3.85 (an AB system, J=16Hz, C—19H₂), 2.0 (s, OAc), 0.53 (s, C—18H₃); IR v_{max} (CHCl_3) 2950 (s), 2900 (sh), 1725 (s), 1420 (m)cm⁻¹; mass spec. molecular ion m/e = 613; (analysis found: % C, 74.18; H, 8.11; N, 6.65; $\text{C}_{38}\text{H}_{51}\text{O}_4\text{N}_3$; requires: % C, 74.35; H, 8.38; N, 6.85).

(b) Ozonolysis
The adduct from (a) above (250 mg) in acetone (10 ml) containing tetracyanoethylene (55 mg, 1 eq) at -78°C was treated with ozone for 3 min (approx. 1.5 eq). The system was purged with argon whilst warming to room temperature. The product mixture was separated by plc to give 130 mg of starting material (nmr) and the corresponding 20(S)-fatty derivative (90 mg, 84%) as a white foam. $^1\text{Hnmr}$ δ 9.55 (d, J=3.75Hz, C—22H), 7.45 (s, 5H, aryl), 5.15 (m, W = 12Hz, C—3H), 4.92 and 4.82 (an AB system, J=10Hz, C—6H, 7H), 4.18 and 4.70 (an AB system J=16Hz, C—19H₂), 2.0 (s, OAc), 1.12 (d, J=7H, C—21H₃), 0.57 (s, C—18H₃).

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Example 2

- (a) Reaction of ergocalciferol acetate with phthalazine-1,4-dion
- Phthalhydrazid (10 g, 2.5 eq) was suspended in a solution of rg calciferolacetat (10 g) in dry CH_2Cl_2 (200 ml). The efficiently mixed mixture was cooled to 0°C, and a solution of lead tetra-acetate (20 g) in dry CH_2Cl_2 (100 ml) and acetic acid (1 ml) was added dropwise. The reaction was monitored by tlc. Upon completion, the residual phthalhydrazide was filtered off. Aqueous work-up followed by careful crystallisation from ethylacetate gave 7.4 g (54%) of 6(R),19-[N,N'-phthalhydrazido]-9,10-seco- β -acetoxy-ergosta-5(10), 7(E), 22(E)-triene. m.p. 202–203°C; $[\alpha]_D = +343^\circ$ ($c = 1.02$); UV λ_{max} 238 nm (38250) and 312nm (11300); $^1\text{Hnmr}$ δ (8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.08 (m, W=10Hz, C—3H, 22H, 23H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 2.0 (s, OAc), 0.13 (s, C—18H₃); IR v_{max} (2950 (s), 2900 (sh), 1750 (s), 1660 (s), 1610 (m), 1380 (m), 1355 (m), 1250 (s)cm⁻¹; mass spec. molecular ion m/e = 598; (analysis found: % C, 75.92; H, 8.30; N, 4.61; $\text{C}_{38}\text{H}_{50}\text{O}_4\text{N}_2$ requires: % C, 76.22; H, 8.42; N, 4.68). The mother liquors were chromatographed on silica gel to give 3.6 g (26%) of essentially pure 6(S),19-[N,N'-phthalhydrazido]-9,10-seco- β -acetoxy-ergosta-5(10), 7(E), 22(E)-triene. Solid from CH_2Cl_2 /hexane. m.p. 114–116°C; $[\alpha]_D = -306^\circ$ ($c = 0.64$); $^1\text{Hnmr}$ δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.12 (m, W=9Hz, C—22H, 23H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.1 (m, C—3H), 0.18 (s, C—18H₃). IR v_{max} 3550 (br), 2950 (s), 2900 (sh), 1650 (s), 1610 (m), 1375 (m), 1350 (m)cm⁻¹; mass spec. molecular ion m/e = 598; (analysis found: % C, 77.76; H, 8.78, N, 5.17; $\text{C}_{36}\text{H}_{48}\text{O}_3\text{N}_2$ requires: % C, 77.66; H, 8.69; N, 5.03).
- (b) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco- β -hydroxy-ergosta-5(10),7(E), 22(E)-triene
To the acetate from (a) above (5 g) in benzene (100 ml) were added NaOH/ CH_3OH (1.25 M solution 12 ml). After 20 min, the mixture was diluted with water and CH_2Cl_2 . Acid work-up gave an essentially quantitative yield (4.5 g) of the title β -hydroxy compound, crystalline from CH_2Cl_2 /ether. m.p. 169–171°C; $[\alpha]_D = +392^\circ$ ($c=0.773$); $^1\text{Hnmr}$ δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.12 (m, W=9Hz, C—22H, 23H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.1 (m, C—3H), 0.18 (s, C—18H₃). IR v_{max} 3550 (br), 2950 (s), 2900 (sh), 1650 (s), 1610 (m), 1375 (m), 1350 (m)cm⁻¹; mass spec. molecular ion m/e = 556; (analysis found: % C, 77.76; H, 8.78, N, 5.17; $\text{C}_{36}\text{H}_{48}\text{O}_3\text{N}_2$ requires: % C, 77.66; H, 8.69; N, 5.03).
- (c) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco- β -tetrahydropyranloxyergosta-5(10), 7(E), 22(E)-triene
The alcohol from (b) above (4.5 g) in benzene (100 ml) was stirred overnight with dihydropyran (10 ml) and p-toluene sulphonic acid (10 mg). Aqueous work-up gave the title THP ether (204c) (5 g, 96%). Crystalline from CH_2Cl_2 /ether. m.p. 151–154°C; $[\alpha]_D = +332^\circ$ ($c = 1.25$); $^1\text{Hnmr}$ δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W = 10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.07 (m, W=9Hz, C—22H, 23H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.7 (m, THP, C—2'H), 4.02 (m, C—3H), 3.5 (m, W=20Hz, THP, C—6'H₂), 0.17 (s, C—18H₃); IR v_{max} 2950 (s), 2900 (sh), 1650 (s), 1610 (m), 1370 (s), 1350 (s), 1030 (s)cm⁻¹; mass spec. molecular ion m/e = 640.
- (d) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco- β -[t-butylidimethylsilyloxy]ergosta-5(10), 7(E), 22(E)-triene
The alcohol from (b) above (4.5 g) in CH_2Cl_2 (20 ml) was treated with t-butyl-dimethylsilylchloride (1.9 g) and imidazole (2.7 g) at room temperature for 1.5 hr. Addition of water followed by acid work-up and crystallization from CH_2Cl_2 /hexane gave 5.1 g (94%) of the silyl ether. m.p. 203–205°C; $[\alpha]_D = +313^\circ$ ($c = 1.5$); $^1\text{Hnmr}$ δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.08 (m, W=9Hz, C—22H, 23H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.03 (m, C—3H), 0.88 (s, t-butyl), 0.17 (s, C—18H₃), 0.07 (s, Si—CH₃), 0.05 (S, Si—CH₃); IR v_{max} 2950 (s), 2900 (sh), 1650 (s), 1610 (m), 1370 (s), 1350 (s), 1090 (s)cm⁻¹; mass spec. molecular ion m/e = 670; (analysis found: % C, 74.98; H, 9.26; N, 4.13; $\text{C}_{42}\text{H}_{62}\text{O}_3\text{N}_2\text{Si}$ requires: % C, 75.18; H, 9.31; N, 4.18).
- (e) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco- β -methoxyethoxymethoxyergosta-5(10), 7(E), 22(E)-triene
The alcohol from (b) above (4.5 g) in CH_2Cl_2 (100 ml) was stirred overnight at room temperature with methoxyethoxymethylchloride (8 ml) in the presence of diisopropylethylamine (20 ml). Acid work-up followed by chromatography and crystallisation from CH_2Cl_2 /hexane gave 4.3 g (83%) of the MEM ether. m.p. 123–125°C; $[\alpha]_D = +325^\circ$ ($c = 1.295$); $^1\text{Hnmr}$ δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.15 (m, W=9Hz, C—22H, 23H), 4.82 (s, —OCH₂O—), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.0 (m, C—3H), 3.67 (m, W=6Hz, —OCH₂CH₂O—), 3.43 (s, OCH₃), 0.18 (s, C—18H₃); IR v_{max} 2950 (s), 2900 (sh), 1650 (s), 1610 (m), 1470 (m), 1450 (m), 1370 (s), 1340 (s), 1100 (s), cm⁻¹; mass spec. molecular ion m/e = 644; (analysis found: % C, 74.72; H, 8.57; N, 4.13; $\text{C}_{40}\text{H}_{56}\text{O}_5\text{N}_2$ requires: % C, 74.50; H, 8.75; N, 4.34).
- (f) General procedure for the ozonolysis of the ergostane side chain
The adduct (from (a), (c), (d) or (e) above (4–5 g) in CH_2Cl_2 (130 ml) and methanol (60 ml) was cooled to –78°C. The efficiently mixed solution was treated with an oz n -oxygen mixtur (approx. 1 mmol O_3/min) for 8–12 min (tlc control) and then thoroughly purged with dry argon for approx. 5 min. Triphenylphosphine (2.5–3 g) was added and the mixture, after approx. 30 min at –78°C (tlc monitoring of the breakdown

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of the methoxyhydroperoxide intermediates) was shaken with 5% aqueous NaHCO₃ (to prevent dimethyl acetal formation) and allowed to warm to room temperature. The layers were separated and the organic solution dried. Chromatography through silica gel (40—50 g) gave the aldehyde (75—86%) free from any of the C—20 (R) epimer (nmr). The following compounds were prepared in this manner.

- 5 1) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β-acetoxy-20(S)-formyl-pregna-5(10),7(E)-diene
 Crystalline from CH₂Cl₂/ether. m.p. 192—193°C; [α]_D = +382° (c = 1.235); ¹Hnmr δ 9.55 (d, J=3Hz, C—22H), 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.17 (m, C—3H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 2.07 (s, OAc), 1.07 (d, J=7Hz, C—21H₃), 0.22 (s, C—18H₃); IR vmax (CHCl₃) 2950 (m), 2900 (sh), 1740 (s), 1645 (s), 1610 (m), 1370 (m), 1350 (m), cm⁻¹; mass spec. molecular ion m/e = 530; (analysis found: % C, 72.13; H, 7.12; N, 5.20; C₃₂H₃₈O₅N₂ requires: % C, 72.43; H, 7.22; N, 5.28).
- 10 2) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β-tetrahydropyranloxy-20(S)-formyl-pregna-5(10), 7(E)-diene
 Crystalline from CH₂Cl₂/ether. m.p. 154—156°C; [α]_D = +356° (c = 0.84); ¹Hnmr δ 9.42 (d, J=3Hz, C—22H), 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.69 (m, THP, C—2'H), 4.0 (m, C—3H), 3.5 (m, W=18Hz, THP, C—6'H₂), 0.95 (d, J=6Hz, C—21H₃), 0.23 (s, C—18H₃). IR vmax 2950 (s), 2900 (sh), 1725 (s), 1640 (s), 1610 (m), 1370 (m), 1350 (m), 1025 (s), cm⁻¹; mass spec. molecular ion m/e = 572; (analysis found: % C, 72.89; H, 7.58; N, 4.78; C₃₅H₄₄O₅N₂ requires: % C, 73.40; H, 7.74; N, 4.89).
- 15 3) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β-[t-butylidimethylsilyloxy]-20(S)-formyl-pregna-5(10), 7(E)-diene
 Crystalline from CH₂Cl₂/hexane. m.p. 195—197°C; [α]_D = +335° (c = 1.64); ¹Hnmr δ 9.52 (d, J=3Hz, C—22H), 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.07 (m, C—3H), 1.07 (d, J=7Hz, C—21H₃), 0.88 (s, t-butyl), 0.22 (s, C—18H₃), 0.07 (s, Si—CH₃), 0.03 (s, Si—CH₃); IR vmax 2950 (s), 2900 (sh), 1740 (s), 1650 (s), 1610 (s), 1350 (s), 1090 (s), cm⁻¹; mass spec. molecular ion m/e = 602; (analysis found: % C, 71.57; H, 8.49; N, 4.51; C₃₆H₅₀O₄N₂Si requires: % C, 71.72; H, 8.36; N, 4.65).
- 20 4) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-20(S)-formyl-3β-methoxyethoxymethoxy-pregna-5(10), 7(E)-diene
 Crystalline from CH₂Cl₂/hexane, m.p. 136—137°C; [α]_D = +327° (c = 0.62); ¹Hnmr δ 9.49 (d, J=3Hz, C—22H), 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 4.87 (s, —OCH₂O—), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.03 (m, C—3H), 3.7 (m, W=6Hz, —OCH₂CH₂O—), 3.47 (s, OCH₃), 1.07 (d, J=7Hz, C—21H₃), 0.22 (s, C—18H₃); IR vmax 2950 (m), 2900 (sh), 1740 (m), 1650 (s), 1610 (m), 1370 (m), 1350 (m), 1030 (m).

Example 3

- 40 General procedure for the reduction of the C—20(S)-formyl to the C—20(S)-(hydroxyethyl) derivative
 The aldehyde compound (2.5—3.5 g) in benzene (60—90 ml) was added dropwise over a 15—20 min period to NaBH₄ (0.8—1.0 g) in ethanol (20—30 ml). After the addition, the excess reducing agent was carefully quenched with dilute aqueous HCl. The mixture was diluted with CH₂Cl₂. Aqueous work-up gave the desired alcohol in essentially quantitative yield. The following compounds have been prepared in this manner.

- 45 1) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β-acetoxy-20(S)-[hydroxymethyl]-pregna-5(10), 7(E)-diene
 Crystalline from CH₂Cl₂/ether. m.p. 238—240°C; [α]_D = +363° (c = 0.875); ¹Hnmr δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.07 (m, C—3H), 4.78 and 4.21 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 3.47 (m, W=14Hz, C—22H₂), 2.05 (s, OAc), 1.0 (broad singlet, C—21H₃), 0.17 (s, C—18H₃); IR vmax (CHCl₃) 3200 (br), 2950 (m), 2900 (sh), 1750 (m), 1650 (s), 1610 (m), 1380 (m), 1350 (m), cm⁻¹; mass spec. molecular ion m/e = 532.
- 50 2) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-20(S)-[hydroxymethyl]3β-tetrahydropyranloxy-pregna-5(10), 7(E)-diene
 Crystalline from CH₂Cl₂/ether, m.p. 170—173°C; [α]_D = +341° (c = 0.58); ¹Hnmr δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.67 (m, THP, C—2'H), 4.0 (m, C—3H), 3.5 (m, W=18Hz, C—22H₂, THP, C—6'H₂), 1.0 (broad singlet, C—21H₃), 0.19 (s, C—18H₃); IR vmax 3600 (br), 2950 (s), 2900 (sh), 1650 (s), 1610 (m), 1370 (m), 1350 (m), 1025 (m), cm⁻¹; mass spec. molecular ion m/e = 574; (analysis found: % C 72.96; H, 7.96; N, 4.73; C₃₅H₄₆O₅N₂ requires: % C, 73.14; H, 8.07; N, 4.87).
- 55 3) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β-[t-butylidimethylsilyloxy]-20(S)-[hydroxymethyl]-pr gna-5(10), 7(E)-diene
 Crystalline from CH₂Cl₂/hexane. m.p. 145—148°C; [α]_D = +312° (c = 1.22); ¹Hnmr δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 4.78 and 4.22 (an AB syst m, J=18Hz, C—19H₂),

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4.75 (d, J=10Hz, C—6H), 4.03 (m, C—3H), 3.4 (m, W=14Hz, C—22H₂), 1.0 (broad singlet, C—21H₃), 0.88 (s, t-butyl), 0.19 (s, C—18H₃), 0.07 (s, Ci—CH₃)₂; IR vmax 3500 (br), 2950 (s), 2900 (sh), 1640 (s), 1610 (m), 1340 (s), 1250 (s), 1090 (s), cm⁻¹; mass spec. molecular ion m/e = 604; (analysis found: % C, 71.56; H, 8.70; N, 4.47; C₃₈H₅₂O₄N₂Si requires: % C, 71.48; H, 8.67; N, 4.63).

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Example 4

6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β-acetoxy-20(S)-ethenylpregna-5(10), 7(E)-diene

Methyltriphenylphosphonium bromide (60 mg, 1.2 eq) was suspended in THF (6 ml). n-Butyl lithium (1.5 M solution 0.15 ml) was added. To the resulting orange-coloured solution, the 3β-acetoxy aldehyde from Example 2(f) (1) (100 mg) in benzene (6 ml) was added quickly. After a further 10 min, water was added and the mixture extracted with CH₂Cl₂. Acid work-up followed by purification by plc gave 75 mg (75%) of the title product. Crystalline from CH₂Cl₂/ether. m.p. 173—175°C; [α]_D = +386° (c = 0.86); ¹Hnmr δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.6—4.8 (m, C—3H, 22H, 23H₂), 4.78 and 4.21 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 2.03 (s, OAc), 0.95 (d, J=7Hz, C—21H₃), 0.17 (s, C—18H₃); IR vmax 2950 (m), 1740 (s), 1650 (s), 1610 (m), 1370 (s), 1350 (s), 1260 (s), 1230 (s), cm⁻¹; mass spec. molecular ion m/e = 528; (analysis found: % C, 75.03; H, 7.72; N, 5.21; C₃₃H₄₀O₄N₂ requires % C, 74.97; H, 7.63; N, 5.30).

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Example 5

(a) Preparation of isobutylene epoxide

To methylallyl chloride (200 ml), 186 g) cooled in an ice bath was added 80% H₂SO₄ (H₂SO₄, 95%, 109 ml; H₂O, 40 ml, 1 eq) over a 30 min period. The temperature of the mixture was maintained between 25 5—10°C. After a further 3 hr the mixture was added to ice and diluted to a total volume of approx. 100 ml. The layers were separated and the organic residue distilled to remove the by-product β,β-dimethylvinyl chloride and unreacted starting material. These materials are removed below 80°C. The darkly coloured distillation residue is 1-chloro-2-methyl-propan-2-ol (128a) δ 3.47 (s, 2H), 2.97 (s, 1H, exchanges with D₂O), 1.32 (s, 6H). This material was used without further purification.

To a 500 ml round bottom flask containing KOH (200 g) in water (125 ml) at 80°C and fitted with a mechanical stirrer and condenser, was added dropwise the crude chlorohydrin. The crude epoxide distilled directly from the reaction mixture. Redistillation gave isobutylene epoxide (48 g, 35%), b.p. 51°C (lit.^{128c} 52°C); ¹Hnmr δ 2.6 (s, 2H), 1.33 (s, 6H).

(b) 4-bromo-2-methyl-2-hydroxy-butane

To ethyl-3-bromo-propionate (21 g) in ether (150 ml) at 0°C was added methylmagnesium bromide (3 M soln. in ether, 125 ml, excess) dropwise. After the addition was complete, the mixture was stirred for a further 2 hrs at room temperature. After cooling again to 0°C, the mixture was carefully quenched with NH₄Cl (30 g) in water (200 ml). The layers were separated and the ether layer washed with water until neutral, followed by brine, and dried. Evaporation gave the crude bromo-alcohol (227a) ¹Hnmr δ 3.53 (t, J=9Hz, 2H), 2.93 (s, 1H, exchanges with D₂O), 2.07 (t, J=9Hz, 2H), 1.27 (s, 6H); [lit.¹⁷¹ 3.54 (t, J=8.5Hz, 2H), 2.65 (s, broad, 1H), 2.10 (t, J=8.5Hz, 2H), 1.26 (s, 6H)].

(c) 4-Bromo-2-methyl-2-(triethylsilyloxy)-butane

Half of the crude bromide from (b) above in ether (50 ml) containing pyridine (5 ml), imidazole (10 g) and triethylsilylchloride (10 ml) was stirred for 2 days at room temperature. Water was added. Acid-work-up followed by chromatography gave 11 g (62% from the propionate) of the desired compound, homogeneous by tlc. ¹Hnmr δ 3.52 (m, 2H), 2.03 (m, 2H), 1.25 (s, 6H), 1.2—0.2 (m, 15H); IR vmax (thin film) 3000 (s), 2950 (sh), 1460 (m), 1420 (m), 1380 (m), 1365 (m), 1230 (s), 1195 (s), 1170 (m), 1100 (s), 1040 (s), 1010 (s), 965 (m), 840 (w), 740 (s), 720 (s), cm⁻¹.

(d) 3-Methyl-2-buten-1-yl-triphenylphosphonium bromide

The bromide from (c) above (1 g) and triphenylphosphine (0.9 g) in benzene (4 ml) was thoroughly degassed, and then heated to reflux. After 3 days, the insoluble material was filtered off to give 1.2 g (85%) of phosphonium salt (228). Recrystallised from CH₂Cl₂/ETOAc. m.p. 234—238°C (lit.^{118b} 236—239°C); ¹Hnmr δ 8.17—7.67 (m, 15H, aryl), 5.18 (m, W=18Hz), 4.73—4.2 (m, 2H), 1.67 (d, J=5Hz, 3H), 1.31 :d, J=5Hz, 3H); IR vmax 2900 (w), 1590 (w), 1490 (m), 1435 (s), 1110 (s), cm⁻¹.

(e) Methyl diphenylphosphine oxide

Methyltriphenylphosphonium bromide (6 g) was refluxed overnight with KOH (5 g) in water (70 ml). The mixture was allowed to cool to room temperature and then extracted (3x) with CH₂Cl₂. The organic layer was washed with brine, dried and the solvent removed to give the crude solid product (3.5 g) in essentially quantitative yield. Recrystallised from acetone. m.p. 113—114°C (lit.¹³² 109—111°C); ¹Hnmr δ 8.0—7.3 (m, 10H, aryl), 2.03 (d, J=13Hz, 3H); IR vmax 1440 (s), 1175 (s), cm⁻¹; mass spec. molecular ion m/z = 216.

(f) 3-Hydroxy-3-methylbut-1-yl-diphenylphosphine oxide

Methyldiphenylphosphine oxide (1.5 g) was suspended in ether (20 ml) at 0°C. BuLi (1.2 eq) was slowly added, and an orange coloured solution formed. To this was added isobutylene epoxide (0.8 ml, 1.3 eq). After approx. 15 min, the mixture was carefully quenched with water. This mixture was extracted with CH₂Cl₂ (2x) and the organic layer was washed with 4% aqueous HCl/brine and concentrated. The resulting yellow, oily product was dissolved in a water-ether mixture and the layers separated. The ether layer was washed once with water, and the combined aqueous fractions extracted with CH₂Cl₂ (3x). The organic layer was washed with brine and dried. The solvent was removed and the resulting colourless oil was taken up in benzene and refluxed through a soxhlet containing CaH₂ for 2 hr. The solvent was removed to give crude title compound (1.4 g, 70%; as an oil. ¹Hnmr δ 8.0—7.3 (m, 10H, aryl), 2.5 (m, W=34Hz, 2H), 1.83 (m, W=32Hz, 3H), 1.23 (s, 6H); IR vmax (CCl₄) 3500 (m), 2950 (m), 1440 (s), cm⁻¹.

(g) 3-tetrahydropyranloxy-3-methylbut-1-yl-diphenylphosphine oxide

The phosphine oxide from (f) above (1.4 g) was dissolved in dihydropyran (20 ml) and benzene (5 ml). p-Toluenesulphonic acid (10 mg) was added. After 20 hr, the mixture was concentrated, added to CH₂Cl₂ and washed with 5% aqueous NaHCO₃/brine and dried. Evaporation of the solvent gave the crude product (1.8 g) essentially quantitatively as a solid. Recrystallised from acetone. m.p. 146—148°C; ¹Hnmr δ 8.0—7.3 (m, 10H, aryl), 4.67 (m, W=6Hz, THP, C—2'H), 3.67 (m, W=36Hz, THP, C—6'H₂), 1.23 (s, 6H); IR vmax 2950 (m), 1440 (m), 118C (s), cm⁻¹; (analysis found: % C, 70.80; H, 7.73; P, 8.54; C₂₂H₂₉O₃P requires % C, 70.96; H, 7.85; P, 8.32.

(h) Preparation of the lithium bromide adduct of the betaine

To methyltriphenylphosphonium bromide (2.898 g) suspended in ether (50 ml) cooled to 0°C was added butyl lithium (2.03 M soln.; 4 ml). Isobutylene epoxide (1.0 ml, 1.25 eq) was added and some insoluble material instantly formed. After stirring for 15 min, the reaction mixture was allowed to settle and the supernatant liquid was removed. The resulting solid was suspended in ether and transferred to two centrifuge tubes, and spun. The ether was removed. This process was repeated until the ether washing were colourless (usually 4x). The colourless solid material was dried to give the Li-Br adduct of the betaine (1.5 g) 42%. Beilstein and lithium ion positive, flame tests. IR vmax (nujol) 3500 (br), 3000 (s), 1440 (s), cm⁻¹.

30

(i) [3-(triethylsilyloxy)-3-methylbut-1-yl]-triphenylphosphonium tetraphenyl borate

To methyltriphenylphosphonium bromide (3 g) suspended in THF (40 ml) was added phenyl lithium (1 eq; 6 ml of a 1.5 M soln.). After 15 min isobutylene epoxide (1 ml, 1.25 eq) was added followed, after a further 5 min, by a second addition of phenyl lithium (1 eq). To this mixture was added benzophenone (1 g; approx. 0.3 eq). After stirring for 20 min, the reaction was quenched with 48% aqueous HBr until just acidic (litmus paper). The organic solvent was removed on a rotary evaporator, water was added and the aqueous layer washed with ether, and the layers separated. The water was removed (rotary evaporator) and the resulting oil taken up in CH₂Cl₂. Aqueous work-up gave the phosphonium salt (226) (3.1 g) 58% as an oil. ¹Hnmr δ 8.17—7.67 (m, 15H, aryl), 5.37 (broad s, —OH), 3.8 (m, W=32Hz C—1H₂), 1.8 (m, W=22Hz, C—2H₂), 1.28 (s, (—CH₃)₂); IR vmax (CHCl₃) 3450 (s), 3000 (s), 1590 (sh), 1440 (s), cm⁻¹.

(j) Silylation

To the phosphonium salt from (i) above (3.7 g) in CH₂Cl₂ (70 ml) was added imidazole (3.4 g) followed by triethylsilylchloride (5 ml). After 40 hr stirring at room temperature, water was added and the mixture diluted with CH₂Cl₂. The CH₂Cl₂ solution after an acid work-up was evaporated and the oily residue partitioned between water and hexane/ether. The water was evaporated and the residue taken up in CH₂Cl₂ which was washed with brine and dried on evaporation the salt (226b) (3.6 g, 77%) as an oil.

(k) Anion exchange

To the phosphonium salt from (j) above (3.6 gg) in 95% ethanol (50 ml) was added dropwise, with stirring, a solution of sodium tetraphenyl borate (2.5 g; 1.1 eq) in water (20 ml). An oily residue is formed which solidifies on continued stirring. Filtration gives the phosphonium tetraphenyl borate salt (4.78 g, 92%) as a white, amorphous, non-hygroscopic solid which may be recrystallised from acetone/hexane/ethanol. m.p. 150—151°C; ¹Hnmr δ (acetone-d₆) 8.2—6.8 (m, 35H, aryl), 3.53 (m, W=34Hz, C—1H₂), 1.8 (m, W=24Hz, C—2H₂), 1.33 (s, (—CH₃)₂), 1.25—0.5 (m, 15H, —SiEt₃); IR vmax 3100 (s), 2950 (s), 1580 (m), 1490 (s), 1440 (s), 1110 (s), 1020 (s), cm⁻¹; (analysis found: % C, 81.41; H, 7.73; P, 3.93; C₅₃H₆₀BOPSi requires: % C, 81.31; H, 7.73; P, 3.96.

(l) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β-acetoxy-25-hydroxy-cholesta-5(10),7(E),22(E)-triene

60 Method A

To methyltriphenylphosphonium bromide (2.898 g) suspended in THF (32 ml) at 0°C was added butyl lithium (2.03 M, 4 ml). Iso-butylene epoxide (720 μl, 1 eq) was slowly added. After a further 15 min, butyl lithium (4 ml) was added. To 3 ml of this solution was added the aldehyde from Example 2(f) (1) (300 mg) in benzene (10 ml). The red colour was quickly discharged. Water was added and the mixture extracted with CH₂Cl₂. After acid work-up the major product was isolated by plc to give the title compound (105 mg, 31%).

Method B

5 The b taine from (h) above (628 mg) was suspended in ether (15 ml) and THF (10 ml). Butyl lithium was added dropwise until a stable colour was formed and then a further amount (0.75 ml, 2 eq for steroid, 1 eq for P compound) was added. To this mixture was added the aldehyde from Example 2(f) (1) (400 mg) in benzene (6 ml) (approx. 5 min). After the addition, water was added and the mixture extracted with CH_2Cl_2 . Work-up as above gave the title compound (155 mg, 34%).

Method C

10 The phosphonium salt from (h) above (280 mg, 1.5 eq) was dissolved in THF (15 ml) at 0°C. Phenyl lithium (3 eq) was added. The aldehyde (206a) (150 mg, 1 eq) in benzene (6 ml) was added quickly. TLC showed no change during 30 min and so water was added. Work-up as above, and isolation by plc gave the title product (80 mg, 47%). Crystalline from CH_2Cl_2 /ether. m.p. 175–177°C; $[\alpha]_D = +347^\circ$ ($c = 0.83$); $^1\text{Hnmr}$ δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.27 (m, W=10Hz, C—3H, 22H, 23H), 4.78 and 4.21 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 2.03 (s, OAc), 1.15 (s, C—26H₃, 27H₃), 0.97 (d, J=7Hz, C—21H₃), 0.17 (s, C—18H₃); IR v_{max} 3800 (m), 2950 (s), 2900 (sh), 1750 (s), 1650 (s), 1610 (m), 1370 (s), 1350 (s), 1240 (s), 965 (m), cm⁻¹; mass spec. molecular ion m/e = 600; (analysis found: % C, 73.94; H, 8.17; N, 4.59; $\text{C}_{37}\text{H}_{48}\text{O}_5\text{N}_2$ requires: % C, 73.97; H, 8.05; N, 4.66).

20 (m) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β-acetoxy-25-hydroxy-cholesta-5(10),7(E),22(Z)-triene

To the phosphonium salt from (i) above (1.9 g) in THF (30 ml) was added phenyl lithium (1.5 M soln., 1.7 ml, 1 eq). After a few minutes, the aldehyde (206a) (1 g) in benzene (35 ml) was added dropwise over about 1 min. After a further 3 min, water was added and the mixture diluted with CH_2Cl_2 and given an acid work-up. The reaction was repeated as above and the combined products chromatographed to yield 2.12 g (78%) of a crude, yellow coloured product.

25 The above mixture (1.4 g) was treated with AcOH:H₂O:THF (8:1:1) (10 ml) for 1.5 hr. Dilution with CH_2Cl_2 followed by aqueous work-up, chromatography and crystallisation gave 1 g of product (85%). Further recrystallisation from CH_2Cl_2 /ether, indicated the major component to have the following characteristics. m.p. 182–184°C; $[\alpha]_D = +339^\circ$ ($c = 0.84$); $^1\text{Hnmr}$ δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.27 (m, W=12Hz, C—3H, 22H, 23H), 4.78 and 4.21 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 2.03 (s, OAc), 1.17 (s,s C—26H₃, 27H₃), 0.9 (d, J=7Hz, C—21H₃), 0.17 (s, C—18H₃); IR v_{max} 3650 (m), 2950 (s), 2900 (sh), 1750 (s), 1650 (s), 1610 (m), 1370 (s), 1350 (s), 1240 (s), cm⁻¹; mass spec. molecular ion m/e = 600; (analysis found: % C, 74.10; H, 8.15; N, 4.47; $\text{C}_{37}\text{H}_{48}\text{O}_5\text{N}_2$ requires: % C, 73.97; H, 8.05; N, 4.66).

35 (n) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β,25-dihydroxy-cholesta-5(10),7(E)-diene

The unsaturated side chain compound from (i) above (450 mg) in benzene (5 ml) and ethanol (5 ml) containing NaHCO₃ (100 mg) and 5% Pt/C (150 mg) was stirred under a hydrogen atmosphere for 24 hr. The mixture was filtered through celite and the solvent removed. To the residue, in benzene (10 ml), was added 40 NaOH in methanol (1.25 M soln, 2 ml) and the mixture stirred for 20 min at room temperature. Acid work-up and crystallisation from CH_2Cl_2 /ether afforded 380 mg (91%) of the title side chain saturated diol. m.p. 174–177°C; $[\alpha]_D = +408^\circ$ ($c = 0.825$); $^1\text{Hnmr}$ δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 4.78 and 4.22 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 4.11 (m, C—3H), 1.22 (s, C—26H₃, 27H₃), 0.87 (broad singlet, C—21H₃), 0.18 (s, C—18H₃); IR v_{max} 3550 (s), 2950 (s), 2900 (sh), 45 1650 (s), 1610 (m), 1370 (s), 1350 (s), cm⁻¹; (analysis found: % C, 74.65; H, 8.66; N, 5.06; $\text{C}_{35}\text{H}_{48}\text{O}_4\text{N}_2$ requires: % C, 74.96; H, 8.63; N, 5.00).

Example 6

50 General procedure for the conversion of the phthalazine-1,4-dione adduct to the corresponding 5(E),7(E),10(19)-triene system of the calciferol

The adduct (200–600 mg) was refluxed overnight, under argon in ethanol (10 ml) and hydrazine (3 ml). After cooling to room temperature, the solvents were removed under reduced pressure and the resulting solid taken up in water (30 ml) and CH_2Cl_2 (30 ml). To this two-phase system under argon was added dianisyltellurium oxide (150–450 mg), K₂CO₃ (6 g) and 1,2-dibromotetrachloroethane (3 g), and the mixture stirred for approx. 5 hr (tlc control). After acid work-up the mixture was chromatographed through silica gel (12 g) and the product removed from traces of tellurium oxidant by plc to give the desired vitamin D compound in 85–93% yield.

60 (1) 9,10-seco-3β,25-dihydroxy-cholesta-5(E),7(E),10(19)-triene

Prepared from the adduct (240b) (200 mg) as described above, to give 131 mg (92%). Solid from ether/hexane. m.p. 79–81°C; $[\alpha]_D = +160^\circ$ ($c = 0.735$); UV λ_{max} 273 nm (21500); $^1\text{Hnmr}$ δ 6.5 and 5.83 (ABq, J=11Hz, C—6H, 7H), 4.97 (s, C—19H), 4.67 (s, C—19H), 3.85 (m, W=14Hz, C—3H), 1.22 (s, C—26H₃, 27H₃), 0.95 (broad singlet, C—21H₃), 0.55 (s, C—18H₃); IR v_{max} 3400 (m), 2950 (s), 1620 (w); mass spec. molecular ion m/e = 400; (analysis found % C, 77.50; H, 10.99; $\text{C}_{27}\text{H}_{44}\text{O}_2$ requires: % C, 80.94; H, 11.07; $\text{C}_{27}\text{H}_{44}\text{O}_2 \cdot \text{H}_2\text{O}$ requires: % C, 77.46; H, 11.075).

(2) 3 β -(3',5'-dinitrobenzoate) ester

The above crud calciferol (1) (125 mg) in pyridine (5 ml) was treated with 3,5-dinitrobenzoyl chloride (85 mg, 1.1 eq). Water was added and the mixture diluted with ether. After acid work-up, the ster was isolated by plc (129 mg, 70%). Crystalline from ether/hexane, m.p. 105–107°C; $[\alpha]_D = +168$ ($c = 0.97$); $^1\text{Hnmr}$ δ 9.13 (m, 3H, aryl), 6.62 and 5.82 (ABq, J=11Hz, C—6H, 7H), 5.3 (m, W=14Hz, C—3H), 5.07 (s, C—19H), 4.77 (s, C—19H), 1.23 (s, C—26H₃, 27H₃), 0.93 (broad singlet, C—21H₃), 0.43 (s, C—18H₃); IR v_{max} 3550 (m), 2950 (s), 2900 (sh), 1750 (s), 1640 (w), 1550 (s), 1350 (s), 1275 (s), cm⁻¹; (analysis found: % C, 68.62; H, 7.85; N, 4.65; $C_{34}H_{46}N_2O_7$ requires: % C, 68.66; H, 7.80; N, 4.71).

10

Example 7

(a) 9,10-seco-3 β ,25-dihydroxy-cholesta-5(Z),7(E),10(19)-triene

A solution of the 5,6-*trans* compound from Example 6(1) (126 mg) in benzene (30 ml) containing triethylamine (2 drops) and anthracene (25 mg) was thoroughly degassed. A Hanovia lamp (number 654A36) was placed such that the outside of the water cooled jacket was 15 cm from the reaction vessel. The mixture was irradiated for 25 min and the title 5,6-cis compound isolated by plc (93 mg, 74%). Crystalline from acetone/water, m.p. 98–100°C (lit.¹⁷⁴ 95–100°C); $[\alpha]_D = +77$ ($c = 0.26$); UV λ_{max} 262 nm (19060); $^1\text{Hnmr}$ δ 6.25 and 6.1 (ABq, J=11Hz, C—6H, 7H), 5.05 (s, C—19H), 4.83 (s, C—19H), 3.9 (m, W=18Hz, C—3H), 1.27 (s, C—26H₃, 27H₃), 0.95 (broad singlet, C—21H₃), 0.55 (s, C—18H₃); IR v_{max} 3500 (s), 2950 (s), 2900 (sh), 1640 (w), 1480 (m), 1380 (m), 1055 (s), cm⁻¹; (analysis: $C_{27}H_{44}O_2 \cdot H_2O$ requires: % C, 77.46; H, 11.08; found: % C, 77.29; H, 11.08). The melting point of an authentic sample supplied by Roussel Uclaf, Inc. (Romainville France) did not depress on mixing.

(b) 3-(3',5'-dinitrobenzoate) ester

Prepared as previously described in Example 6(2). Crystalline from ether/hexane, m.p. 149–150°C (lit.¹⁷² 147–148°C); $[\alpha]_D = +90$ ($c = 0.6$); (analysis: $C_{34}H_{46}N_2O_7$ requires: % C, 68.66; H, 7.80; N, 4.71; found: % C, 68.94; H, 7.80; N, 4.52).

Example 8

 SO_2 adducts from 9,10-seco-3 β -hydroxy-ergosta-5(Z),7(E),10(19),22(E)-tetraene

Sulphur dioxide was slowly passed through a well-stirred mixture of benzene (100 ml) and water (50 ml) containing ergocalciferol (5 g), for a total of 3.5 hr. After this time, air was passed through the mixture for approx. 20 min. Ether and brine were added and the layers separated. Aqueous work-up gave the known sulphur dioxide adducts (172a, 173a) which were used without further purification.

35

Example 9

(a) 9,10-seco-3 β -(triethylsilyloxy)-ergosta-5(E),7(E),10(19),22(E)-tetraene

To the 3 β -alcohol corresponding to the title compound (4.3 g) in CH_2Cl_2 (50 ml) was added imidazole (4 g) followed by triethylsilylchloride (3 ml). After a few minutes, water was added and the organic layer washed with water/brine and dried. The required silyl ether was isolated essentially quantitatively after chromatography as an oil. UV λ_{max} 274 nm; $^1\text{Hnmr}$ δ 6.45 and 5.87 (ABq, J=11Hz, C—6H, 7H), 5.2 (m, W=9Hz, C—22H, 23H), 4.92 (s, C—19H), 4.63 (s, C—19H), 3.82 (m, W=18Hz, C—3H).

(b) 9,10-seco-1 α -hydroxy-3 β -(triethylsilyloxy)-ergosta-5(E),7(E),10(19),22(E)-tetraene

N-Methylmorpholine N-oxide (NMO) (6.3 g) was stirred with anhydrous $MgSO_4$ in CH_2Cl_2 (50 ml) for 30 min. Selenium dioxide (1.3 g) was stirred in methanol (50 ml) for 45 min and warmed to reflux. The above CH_2Cl_2 mixture was filtered into a solution of the 5,6-*trans*-ergocalciferol derivative from (a) above (5.5 g) in 1,2-dichloroethane (50 ml). This mixture was warmed to reflux and then the hot methanol mixture added, and refluxing of the whole continued for a further 35 min. The heat source was removed and the mixture diluted with CH_2Cl_2 . Aqueous work-up followed by chromatography through silica gel (40 g) gave 2.66 g (47%) of the title compound as an oily product. UV λ_{max} 274 nm; $^1\text{Hnmr}$ δ 6.57 and 5.90 (ABq, J=11Hz, C—6H, 7H), 5.25 (m, W=9Hz, C—22, 23H), 5.08 (s, C—19H), 4.98 (s, C—19H), 4.65–3.92 (m, C—1H, 3H).

(c) 9,10-seco-1 α ,3 β -dihydroxy-ergosta-5(E),7(E),10(19),22(E)-tetraene

The silylether from (b) above (460 mg) in THF (10 ml) was stirred for 30 min with tetrabutylammonium fluoride (460 mg). The mixture was diluted with CH_2Cl_2 and after aqueous work-up, the title diol was purified by plc to give 305 mg (84%). Crystalline from ether/hexane, m.p. 103–105°C; $[\alpha]_D = +172$ ($c = 0.58$); UV λ_{max} 272 nm (22600); $^1\text{Hnmr}$ δ 6.38 and 5.82 (ABq, J=11Hz, C—6H, 7H), 5.18 (m, W=9Hz, C—22H, 23H), 4.9 (m, W=9Hz, C—19H), 4.53–3.77 (m, C—1H, 3H), 0.57 (s, C—18H₃); IR v_{max} 3500 (s), 2950 (s), 2900 (sh), 1640 (w), 1460 (m), 1375 (m), 1050 (s), 1030 (s), cm⁻¹; mass spec. molecular ion m/e = 412; (analysis found: % C, 79.57; H, 10.71; $C_{28}H_{44}O_2 \cdot \frac{1}{2}H_2O$ requires: % C, 79.76; H, 10.76).

(d) 9,10-s co-1 α -hydr xy-3 β -triethylsilyloxy-erg sta-5(Z),7(E),10(19),22(E)-t tra- ne

The 5,6-*trans* compound from (b) above (600 mg) in benzene (30 ml) containing phenazin (120 mg) and triethylamine (few drops) was photoisomerised as above for 30 min to give 400 mg (66%) of the titl

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5,6-cis vitamin. UV λ_{max} 263 nm; $^1\text{Hnmr}$ δ 6.38 and 6.08 (ABq, J=11Hz, C—6H, 7H), 5.23 (m, W=10Hz, C—19H, 22H, 23H), 5.0 (s, C—19H), 4.6—3.92 (m, C—1H, 3H).

() 9,10-seco-1 α -3 β -dihydroxy-ergosta-5(Z),7(E),10(19),22(E)-tetra-ene

The silyl ether derivative from (d) above (200 mg) was stirred at room temperature in THF (10 ml) with N-Bu₄NF (1 M soln. in THF, 2 ml) for about 30 min. Dilution with CH₂Cl₂ and aqueous work-up followed by purification by plc gave 129 mg (82%). Crystalline from ether/hexane gave the title compound. m.p. 141—143°C (lit.¹⁰⁶ 138—140°C); [α]_D = +34° (c = 0.645); UV λ_{max} 264 nm (19100); $^1\text{Hnmr}$ δ 6.35 and 6.05 (ABq, J=11Hz, C—6H, 7H), 5.16 (m, W=14Hz, C—19H, 22H, 23H), 4.98 (s, C—19H), 4.6—3.85 (m, C—1H, 3H), 0.55 (s, C—18H₃); IR v_{max} 3500 (s), 2950 (s), 2900 (sh), 1640 (w), 1460 (m), 1370 (m), 1060 (s), cm⁻¹; mass spec. molecular ion m/e = 412; (analysis: C₂₈H₄₄O₂ requires: % C, 81.50; H, 10.75; O, 7.76; found: % C, 81.39; H, 10.60).

Example 10

(a) 9,10-seco-3 β -acetoxy-1 α -benzoyloxy-ergosta-5(E),7(E),10(19),22(E)-tetra-ene

The 1 α -hydroxy-3 β -triethylsilyloxy-compound from Example 9(b) (2 g) was treated with benzoyl chloride (2 ml) in pyridine (25 ml). After 30 min water was added and the mixture diluted with ether. After acid work-up, the solvent was removed and the resulting oil stirred overnight in THF:H₂O:AcOH; 8:1:3 (36 ml). After dilution with ether and aqueous work-up, the crude benzoate-alcohol was taken up in benzene (40 ml). Triethylamine (7 ml), acetic anhydride (3 ml) and 4-dimethylaminopyridine (15 mg) were added. After 30 min, water was added and the mixture diluted with ether. Acid work-up and chromatography through silica (10 g) gave 1.76 g (83%) of the title acetate-benzoate as an oil. $^1\text{Hnmr}$ δ 8.05 (m, W=12Hz, 2H, aryl), 7.5 (m, W=10Hz, 3H, aryl), 6.58 (d, J=11Hz, C—6H), 5.88 (m, W=16Hz, C—1H, 7H), 5.15 (m, W=10Hz, C—3H, 19H₂, 22H, 23H), 2.05 (s, OAc), 0.57 (s, C—18H₃).

(b) 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-1 α -benzoyloxy-3 β -acetoxy-ergosta-5(10),7(E),22(E)-triene

To a well-stirred suspension of phthalhydrazide (2 g) in CH₂Cl₂ (200 ml) at 0°C, containing the vitamin from (a) above (2 g), was added dropwise a solution of Pb(OAc)₄ (4 g) in CH₂Cl₂ (20 ml) and acetic acid (1 ml). After consumption of starting material (tlc control), the excess phthalhydrazide was removed by filtration. Aqueous work-up and chromatography gave 1.4 g (52% from 248b) of a 95:5 mixture of (250) and the presumed 6(S) isomer. Crystallisation from CH₂Cl₂/hexane gave pure title compound. m.p. 211—213°C; [α]_D = +295° (c = 0.83); $^1\text{Hnmr}$ δ 8.5—7.3 (m, 9H, aryl), 5.95 (m, W=14Hz, C—1H, 7H), 5.28 (m, C—3H), 5.17 (m, W=10Hz, C—22H, 23H), 4.92 and 4.37 (an AB system, J=18Hz, C—19H₂), 4.8 (m, C—6H), 2.05 (s, OAc), 0.17 (s, C—18H₃); IR v_{max} 2950 (s), 2900 (sh), 1750 (s), 1720 (s), 1640 (s), 1610 (m), 1265 (s), 1245 (s), cm⁻¹; mass spec. molecular ion m/e = 718; (analysis found % C, 75.26; H, 7.54; N, 3.82; C₄₅H₅₄O₆N₂; requires: %C 75.18; H, 7.57; N, 3.90).

Example 11

(a) SO₂ adducts of 9,10-seco-3 β -(t-butyldimethylsilyloxy)-ergosta-5(E),7(E),10(19),22(E)-tetraene

The crude mixture of sulphur dioxide adducts of ergocalciferol (prepared from 5 g of ergocalciferol as described previously), in CH₂Cl₂ (40 ml), containing imidazole (4 g) was stirred with t-butyldimethylsilyl chloride (3.5 g). After 1.5 hr, the reaction was worked-up as described previously to give, after chromatography, 4.8 g (66%, from ergocalciferol) of the title compound as an oil epimeric at C—6. $^1\text{Hnmr}$ δ 5.22 (m, W=9Hz, C—22H, 23H), 4.64 (m, W=10Hz, C—6H, 7H), 4.02 (m, W=16Hz, C—3H), 3.67 (broad s, C—19H₂), 0.91 (s, t-Bu), 0.68 + 0.59 (2 × s, C—18H₃), 0.07 s, [(Si—CH₃)₂].

(b) SO₂ adducts of 9,10-seco-3 β -triethylsilyloxy-ergosta-5(E),7(E),10(19),22(E)-tetra-ene

The crude mixture of sulphur dioxide adducts of ergocalciferol (prepared from 5 g of ergocalciferol as described previously), in CH₂Cl₂ (40 ml), containing imidazole (4 g) was stirred with triethylsilylchloride (3.5 ml). After about 30 min, the reaction was worked up as described previously to give, after chromatography, 5.3 g (74% from ergocalciferol) of (210b) as an oil. $^1\text{Hnmr}$ δ 5.22 (m, W=9Hz, C—22H, 23H), 4.64 (m, W=10Hz, C—6H, 7H), 4.02 (m, W=16Hz, C—3H), 3.67 (broad s, C—19H₂).

(c) SO₂ adducts of 9,10-seco-3 β -(t-butyldimethylsilyloxy)-20(S)-formyl-pregna-5(E),7(E),10(19)-triene

The vitamin D₂ adduct from (b) above (4.7 g) was treated with ozone as described in the general procedure to give, after chromatography, 3.25 g (78%) of the aldehyde (211a). $^1\text{Hnmr}$ δ 9.39 (m, C—22H), 4.66 (m, W=16Hz, C—6H, 7H), 4.0 (m, W=16Hz, C—3H), 3.66 (broad s, C—19H₂), 1.15 (d, W=6Hz, C—21H₃), 0.89 (s, t-Bu), 0.71 + 0.62 (2 × s, C—18H₃), 0.05 s, [(Si—CH₃)₂]; IR v_{max} (thin film) 2950 (s), 2900 (sh), 1720 (s), 1660 (w), 1460 (m), 1305 (s), 1250 (s), 1150 (m), cm⁻¹.

(d) Similarly prepared in 82% yield from (b) above was the SO₂ adducts of 9,10-seco-3 β -(triethylsilyl xy)-20(S)-formyl-pregna-5(E),7(E),10(19)-triene. $^1\text{Hnmr}$ δ 9.57 (m, C—22H), 4.67 (m, W=12Hz, C—6H, 7H), 3.97 (m, W=16Hz, C—3H), 3.65 (broad s, C—19H₂), 1.15 (d, J=6Hz, C—21H₃); IR v_{max} (thin film) 2950 (s), 2900 (sh), 1735 (s), 1660 (w), 1460 (m), 1380 (m), 1310 (s), 1150 (m), cm⁻¹.

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Example 12

(a) SO₂ adducts of 9,10-seco-3β-(t-butyldimethylsilyloxy)-20(S)-(hydroxymethyl)-pregna-5(E),7(E),10(19)-triene

The aldehyd corresponding to the title compound (3.1 g) was reduced as described in the general procedure to the title compound in essentially quantitative yield, ¹Hnmr δ 4.63 (m, W=12Hz, C—6H, 7H), 4.02 (m, W=16Hz, C—3H), 3.80—3.28 (m, C—19H₂, 22H₂), 1.05 (d, J=6Hz, C—21H₃), 0.87 (s, t-Bu), 0.68 + 0.58 ([2 × s, C—18H₃], 0.05 [s, (Si—CH₃)₂]; IR max (thin film) 3550 (br), 2950 (s), 2900 (sh), 1660 (w), 1475 (m), 1350 (s), 1275 (s), 1155 (m), cm⁻¹.

10 (1) Similarly prepared in greater than 90% yield was the SO₂ adducts of 9,10-seco-3β-(triethylsilyloxy)-20(S)-(hydroxymethyl)-pregna-5(E),7(E),10(19)-triene, ¹Hnmr δ 4.63 (m, W=12Hz, C—6H, 7H), 3.93 (m, W=16Hz, C—3H), 3.77—3.17 (m, C—19H₂, 22H₂); IR v_{max} (thin film) 3550 (br), 2950 (s), 2900 (sh), 1660 (w), 1460 (m), 1380 (m), 1305 (s), 1240 (m), 1155 (m), cm⁻¹.

15 (2) 9,10-seco-3β-(t-butyldimethylsilyloxy)-20(S)-(hydroxymethyl)-pregna-5(E),7(E),10(19)-triene Adducts of (1) above (3 g) was stirred in refluxing methanol (50 ml) containing NaHCO₃ (3 g) for 2.5 hr. Work-up as described above gave 2.36 g (90%) of the calciferol. UV λ_{max} 274 nm; ¹Hnmr δ 6.47 and 5.87 (ABq, J=11Hz, C—6H, 7H), 4.92 (s, C—19H), 4.65 (s, C—19H), 4.1—3.15 (m, C—3H, 22H₂), 1.06 (d, J=5Hz, C—21H₃), 0.9 (s, t-Bu), 0.58 (s, C—18H₃), 0.07 s, [(Si—CH₃)₂].

20 (3) Similarly prepared in 47% yield from the adducts of (1) above after chromatography was 9,10-seco-3β-(triethylsilyloxy)-2-(S)-(hydroxymethyl)-pregna-5(E),7(E),10(19)-triene (267d). UV λ_{max} 273 nm; ¹Hnmr δ 6.43 and 5.7 (ABq, J=11Hz, C—6H, 7H), 4.9 (s, C—19H), 4.6 (s, C—19H), 4.03—3.13 (m, C—3H, 22H₂).

25 Example 13

9,10-seco-3β-hydroxy-20(S)-hydroxymethyl-pregna-5(E),7(E),10(19)-triene

Method A

The phthalazine adduct from Example 3(1) (200 mg) was treated with hydrazine, followed by oxidation as described in the general procedure to give the title product (105 mg; 85%).

30

Method B

The phthalazine adduct from Example 3(3) (250 mg) was similarly converted to give the t-butyldimethylsilyl ether of the title product (166 mg, 90%). This material in refluxing THF (10 ml) was stirred with n-Bu₄NF (1 M soln. in THF, 2 ml) for 1 hr. Dilution with CH₂Cl₂, followed by aqueous work-up and purification by plc gave (267c) (107 mg, 87%).

Method C

40 The product of Method B (160 mg) obtained via the corresponding SO₂ adducts was similarly converted to the title compound.

Method D

45 The triethylsilyl ether of the title compound (160 mg) obtained via the corresponding SO₂ adducts in THF (10 ml) was stirred at room temperature with n-Bu₄NF (1 M soln. in THF, 2 ml). After about 30 min, the reaction was worked up as for (B) above, to give (267c) (101 mg, 85%).

Crystalline from CH₂Cl₂/hexane. m.p. 104—106°C; [α]_D = +190° (C = 0.37); UV λ_{max} 273 nm (22640); ¹Hnmr δ 6.5 and 5.83 (ABq, J=11Hz, C—6H, 7H), 4.93 (s, C—19H), 4.62 (s, C—19H), 4.08—3.12 (m, C—3H, 22H₂), 1.05 (d, J=5Hz, C—21H₃), 0.58 (s, C—18H₃); IR v_{max} 3450 (s), 2980 (s), 2950 (sh), 1635 (w), 1450 (m), 1050 (s), 1030 (s), cm⁻¹; mass spec. molecular ion m/e = 330; (analysis found: % C, 79.46; H, 9.94; C₂₂H₃₄O₂ requires % C, 79.95; H, 10.37).

50

Example 14

55 6(R),19-[N,N'-phthalhydrazido]-9,10-seco-3β-acetoxy-20(S)-[p-toluenesulphonyloxymethyl]-pregna-5(10),7(E)-diene

The alcohol from Example 3(1) (2.275 g) in pyridine was stirred overnight with p-toluenesulphonyl-chloride (6.25 g) at room temperature. Water was added to the ice cooled mixture and after about 20 min, the mixture extracted with CH₂Cl₂. Acid work-up followed by crystallisation from CH₂Cl₂/ether gave 2.5 g (85%) of the required tosylate (216). m.p. 91—92°C; [α]_D = +308° (c = 1.26); ¹Hnmr δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl; and d, J=7Hz, 2H, tosyl), 7.33 (d, J=7Hz, 2H, tosyl), 5.85 (d, J=10Hz, C—7H), 5.06 (m, C—3H), 4.78 and 4.2 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 3.8 (m, W=12Hz, C—22H₂), 2.43 (s, t syl), 2.03 (s, OAc), 0.88 (d, J=5Hz, C—21H₃), 0.13 (s, C—18H₃); IR v_{max} 2950 (s), 2900 (sh), 1750 (s), 1650 (s), 1610 (s), 1475 (s), 1240 (s), 1175 (s), cm⁻¹; mass spec. molecular ion m/e = 686; (analysis found: % C, 68.09; H, 6.844; N, 4.00; S, 4.90; C₃₉H₄₈O₇N₂S requires: % C, 68.19; H, 6.75; N, 4.08; S, 4.67).

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Example 15

3-m thyl-1-butyn-3-yl tetrahydropyranyl eth r

5 3-Methyl-1-butyl-3-ol (25 ml, 21.7 g), dihydropyran (50 ml) and p-toluen sulphonic acid (5 mg) were mix d tog ther at 0°C for 1 hr, and then stirr d at room temperature for a further 40 hr. The mixture was conc ntrated and the r sidu added to 5% aqueous NaHCO₃ and extracted with benzene. The organic solution was dried to give after distillation 37.3 g (86%) of the title ether. b.p. 47°C/0.8 mm Hg (lit. 30—33°C/0.5 mm⁵⁰; 57°C/3.5 mm¹⁷⁰); ¹Hnmr δ 5.6 (m, THP C2'H), 2.45 (s, C—1H), 1.51 (s, CH₃), 1.48 (s, CH₃); IR .(thin film) 3350 (s), 2950 (s), 2900 (sh), 1125 (s), 1070 (s), 1030 (s), cm⁻¹.

10

Example 16

1-mercaptop-2-methyl-2-hydroxy-propane

15 Ethyl-2-mercaptop acetate (10 ml) was added to dry ether (150 ml). The well-stirred solution was cooled to 0°C, and an ethereal solution of methyl magnesium bromide (3 M soln., 100 ml., 3.3 eq) was added dropwise over 1.5 hr. The mixture was removed from the ice bath and stirred for an additional 30 min. Ammonium chloride (18 g) in water was carefully added, and the mixture neutralised with hydrochloric acid to form 2 clear layers. The layers were separated and the ether layer washed with water/brine and dried. The solvent was removed under reduced pressure and the product distilled to give 4.4 g of the thiol. b.p. 46°C/16 mm Hg (lit. 64°/26 mm^{158a}, 61°/22 mm^{158b}); ¹Hnmr δ 2.6 (d, J=9Hz, C—1H₂), 2.5 (s, exchanges with D₂O, —OH), 1.38 (t, J=9Hz, —SH), 1.28 (s, 6H, —(CH₃)₂); mass spec. m/e 59 (100), 73 (24), 91 (14).

20

Example 17

6(R),19-[N,N'-phthalhydrazido]-23-thia-9,10-seco-3β-acetoxy-25-hydroxy-cholesta-5(10),7(E)-diene

25 To the tosylate from Example 14 (2.51 g) in THF (125 ml) and HMPTA (3 ml) was added 1-mercaptop-2-methylpropan-2-ol from Example 16 (3 ml). The mixture was degassed and NaH (50% dispersion in oil, 1.3 g) was added. After 2 hr, water was added and the mixture diluted with benzene/CH₂Cl₂. Acid work-up followed by chromatography and crystallisation from CH₂Cl₂/ether gave 1.71 g (77%) of the title sulphide. m.p. 187—188°C; [α]_D = +348° (c = 0.62); ¹Hnmr δ 8.3 (m, W=12Hz, 2H, aryl), 7.8 (m, W=10Hz, 2H, aryl), 5.9 (d, J=10Hz, C—7H), 5.07 (m, C—3H), 4.78 and 4.18 (an AB system, J=18Hz, C—19H₂), 4.75 (d, J=10Hz, C—6H), 2.58 (s, C—24H₂), 2.03 (s, OAc), 1.23 (s, C—26H₃, 27H₃), 0.98 (d, J=6Hz, C—21H₃), 0.15 (s, C—18H₃); IR v_{max} 3600 (m), 2950 (s), 2900 (sh), 1740 (s), 1640 (s), 1610 (m), 1370 (s), 1350 (s), 1240 (s), cm⁻¹; mass spec. molecular ion m/e = 620; (analysis found: % C, 69.47; H, 7.63; N, 4.43; S, 5.21; C₃₆H₄₈O₅N₂S requires: % C, 69.64; H, 7.79; N, 4.51; S, 5.17).

30

Example 18

23-thia-9,10-seco-3β,25-dihydroxy-cholesta-5(E),7(E),10(19)-triene

35 Prepared from the adduct of Example 17 as described in the general procedure as an oil. UV λ_{max} 273 nm; ¹Hnmr δ 6.52 and 5.83 (ABq, J=11Hz, C—6H, 7H), 4.95 (s, C—19H), 4.67 (s, C—19H), 3.85 (m, W=14Hz, C—3H), 2.63 (s, C—24H₂), 1.3 (s, C—26H₃, 27H₃), 1.1 (d, W=6Hz, C—21H₃), 0.58 (s, C—18H₃).

40

Example 19

3-(3',5'-dinitrobenzoate)-ester

45 Prepared as described for the compound of Example 6(2) in 67% yield from the adduct of Example 17. Crystalline from ether/hexane. m.p. 108—110°C; [α]_D = +188° (c = 0.742); UV λ_{max} 272 nm (25400) and 262 nm (25400); ¹Hnmr δ 9.12 (m, 3H, aryl), 6.62 and 5.82 (ABq, J=11Hz, C—6H, 7H), 5.33 (m, W=12Hz, C—3H), 5.00 (s, C—19H), 4.78 (s, C—19H), 2.63 (s, C—24H₂), 1.27 (s, C—26H₃, 27H₃), 1.08 (d, W=6Hz, C—21H₃), 0.45 (s, C—18H₃); IR v_{max} 3600 (m), 2950 (s), 2900 (sh), 1740 (s), 1640 (m), 1550 (s), 1350 (s), 1280 (s), 1170 (s), cm⁻¹; mass spec. molecular ion m/e = 612; (analysis found: % C, 64.39; H, 7.26; N, 4.43; S, 5.11; C₃₃H₄₄O₇N₂S requires: % C, 64.68; H, 7.24; N, 4.57; S, 5.23).

50

Example 20

23-thia-9,10-seco-1α-hydroxy-3β,25-bis(triethylsilyloxy)-cholesta-5(E),7(E),10(19)-triene

55 To the diol of Example 18 (400 mg) in CH₂Cl₂ (15 ml) was added imidazole followed by triethylsilyl chloride (450 µl). After 7 hrs, water was added and the mixture diluted with CH₂Cl₂. Acid work-up gave the crude bis TES derivative which was used without further purification.

60 Selenium dioxide (106 mg) was stirred in methanol (5 ml) for 45 min. N-methylmorpholine-N-oxide (NMO) (528 mg) was stirred in CH₂Cl₂ (5 ml) in the presence of anhydrous MgSO₄ for 30 min. The NMO solution was filt r d into a s lution of the crude bis TES derivative in 1,2-dichloro ethane (5 ml) and the mixture warmed to reflux. To this refluxing mixture was add d the SeO₂/methanol. After 35 min at reflux, the h ating mantle was removed, the mixture diluted with CH₂Cl₂ and washed immediately with 5% aqu ous NaHCO₃ and dri d. Purification by plc gave 233 mg [35% from the adduct of Example 17 of the title 1α-hydroxy compound as an oil. UV λ_{max} 274 nm; ¹Hnmr δ 6.58 and 5.92 (ABq, J=12Hz, C—6H, 7H), 5.08 (s, C—19H), 4.97 (s, C—19H), 4.67—4.03 (m, C—1H, 3H), 2.58 (s, C—24H₂), 1.32 (s, C—26H₃, 27H₃).

65

Example 21**23-thia-9,10-seco-1 α ,3 β ,25-trihydroxy-cholesta-5(E),7(E),10(19)-triene**

To the bis TES derivative from Example 20 (112 mg) in THF (5 ml), was added anhydrous tetrabutylammonium fluoride (220 mg) in benzene (3 ml). After 2.25 hr at reflux, the mixture was diluted with ethyl-acetate, washed with water (3×)/brine and dried. The title triol (50 mg, 68%), was isolated by plc. Crystalline from CH_2Cl_2 /hexane. m.p. 129–131°C; $[\alpha]_D = +184^\circ$ ($c = 0.2175$); UV λ_{max} 273 nm (21860); $^1\text{Hnmr}$ δ 6.58 and 5.92 (ABq, J=11Hz, C—6H, 7H), 5.12 (s, C—19H), 5.0 (s, C—19H), 4.65–4.0 (m, C—1H, 3H), 2.67 (s, C—24H₂), 1.28 (s, C—26H₂, 27H₃), 1.12 (d, J=7Hz, C—21H₃), 0.57 (s, C—18H₃); IR ν_{max} 3550 (s), 2950 (s), 2900 (sh), 1640 (w), 1050 (m), 1030 (m), cm⁻¹; mass spec. molecular ion m/e = 434; (analysis found: % C, 71.57; H, 9.57; S, 7.23; $\text{C}_{26}\text{H}_{42}\text{O}_3\text{S}$ requires: % C, 71.84; H, 9.74; S, 7.38).

Example 22**23-thia-9,10-seco-3 β ,25-dihydroxy-cholesta-5(Z),7(E),10(19)-triene**

A solution of the 5,6-trans-trans vitamin from Example 20 (64 mg) in benzene (30 ml) containing triethylamine (1 drop) and antracene (15 mg) was thoroughly degassed and photoisomerised as described in Example 6(1). The mixture was irradiated for 20 min and the required title vitamin (49 mg, 77%) isolated by plc as an oil. UV λ_{max} 262 nm; $^1\text{Hnmr}$ δ 6.25 and 6.0 (ABq, J=11Hz, C—6H, 7H), 5.03 (s, C—19H), 4.82 (s, C—19H), 3.93 (m, W=18Hz, C—3H), 2.67 (s, C—24H₂), 1.27 (s, C—26H₃, 27H₃), 1.08 (d, W=6Hz, C—21H₃), 0.55 (s, C—18H₃).

Example 23

The 3-(3',5'-dinitrobenzoate ester of the product of Example 22 was prepared using the method of Example 6(2). Crystalline from ether/hexane. m.p. 145–148°C; $[\alpha]_D = +109^\circ$ ($C = 0.571$); UV λ_{max} shoulders at 260 nm (24900) and 235 nm (30600); $^1\text{Hnmr}$ δ 9.08 (m, 3H, aryl), 6.33 and 6.06 (ABq, J=11Hz, C—6H, 7H), 5.33 (m, C—3H), 5.15 (s, C—19H), 4.93 (s, C—19H), 2.65 (s, C—24H₂), 1.27 (s, C—26H₃, 27H₃), 1.08 (d, J=6Hz, C—21H₃), 0.57 (s, C—18H₃). IR ν_{max} 3750 (m), 2950 (s), 2900 (sh), 1750 (s), 1640 (s), 1550 (s), 1345 (s), 1280 (s), 1170 (s), cm⁻¹; mass spec. molecular ion m/e = 612; (analysis found: % C, 64.7; H, 7.24; O, 18.25; N, 4.36; S, 5.15; $\text{C}_{33}\text{H}_{44}\text{O}_7\text{N}_2\text{S}$ requires: % C, 64.68; H, 7.24; O, 18.28; N, 4.57; S, 5.23).

Example 24**23-thia-9,10-seco-1 α -hydroxy-3 β ,25-bis(triethylsilyloxy)-cholesta-5(Z),7(E),10(19)-triene**

The corresponding 5(E) compound of Example 20 (180 mg) in benzene (35 ml) containing phenazine (40 mg) and triethylamine (4 drops) was thoroughly degassed and irradiated as described above for 35 min. 137 mg (75%) of the less polar 5(Z) compound was isolated as an oil by plc. UV λ_{max} 263 nm; $^1\text{Hnmr}$ δ 6.35 and 6.05 (ABq, J=11Hz, C—6H, 7H), 5.27 (s, C—19H), 4.95 (s, C—19H), 4.6–3.93 (m, C—1H, 3H), 2.57 (s, C—24H₂), 1.2 (s, C—26H₃, 27H₃).

Example 25**23-thia-9,10-seco-1 α ,3 β ,25-trihydroxy-cholesta-5(Z),7(E),10(19)-triene**

To the corresponding bis TES derivative from Example 24 (185 mg) in THF (8 ml) was added tetrabutylammonium fluoride (1 M soln. in THF, 2 ml). After 1.25 hr at reflux, the mixture was diluted with CH_2Cl_2 . Aqueous work-up and purification by plc gave 110 mg (90%) of the title triol. Crystalline from ether/hexane. m.p. 124–126°C; $[\alpha]_D = +54^\circ$ ($c = 0.37$); UV λ_{max} 264 nm (17400); $^1\text{Hnmr}$ δ 6.35 and 6.05 (ABq, J=11Hz, C—6H, 7H), 5.33 (s, C—19H), 5.0 (s, C—19H), 4.65–4.0 (m, C—1H, 3H), 2.63 (s, C—24H₃), 1.27 (s, C—26H₃, 27H₃), 1.1 (d, J=6Hz, C—21H₃), 0.55 (s, C—18H₃); IR ν_{max} 3550 (s), 2950 (s), 2900 (sh), 1640 (w), 1050 (m), 1030 (m), cm⁻¹; mass spec. molecular ion m/e = 434; (analysis found: % C, 71.63; H, 9.61; S, 7.34; $\text{C}_{26}\text{H}_{42}\text{O}_3\text{S}$ requires: % C, 71.84; H, 9.74; S, 7.38).

Example 26**23-thia-9,10-seco-1 α -bis(3',5'-dinitrobenzoyloxy)-25-hydroxy-cholesta-5(Z),7(E),10(19)-triene**

To the triol from Example 25 (75 mg) in pyridine (3 ml) and benzene (5 ml) was added 3,5-dinitrobenzoylchloride (85 mg). Water was added and the mixture diluted with ether. Work-up as in Example 6(2) and purification by plc gave 97 mg (68%) of the unstable bis (dinitrobenzoate). $^1\text{Hnmr}$ δ 9.8 (m, 6H, aryl), 6.62 (d, J=11Hz, C—6H), 6.12–5.42 (m, C—1H, 3H, 7H, 19H), 5.32 (s, C—19H), 2.63 (s, C—24H₂), 1.27 (s, C—26H₃, 27H₃), 1.08 (broad singlet, C—21H₃), 0.22 (s, C—18H₃).

Example 27**23-thia-9,10-seco-1 α ,3 β ,25-trihydroxy-cholesta-5(Z),7(E),10(19)-triene-23,S-oxides**

The sulphide from Example 25 (100 mg) in methanol (10 ml), ether and water (2 ml) was stirred at room temperature with sodium metaperiodate (50 mg). After 3 hr, a further addition of oxidant (20 mg) was made. After a total of 5 hr, the mixture was diluted with CH_2Cl_2 . Aqueous work-up followed by plc gave 92 mg (89%) of the title sulphoxide mixture. Solid from aceton, methanol/hexane, ether. m.p. 148–155°C; $[\alpha]_D = +77^\circ$ ($c = 0.691$); UV λ_{max} 263 nm (18150) $^1\text{Hnmr}$ δ 6.35 and 6.05 (ABq, J=11Hz, C—6H, 7H), 5.33 (s, C—19H), 5.0 (s, C—19H), 4.62–3.72 (m, C—1H, 3H, —OH, exchanges with D_2O), 2.83 and 2.75 (br ad singlets, C—24H₂), 1.52 and 1.38 (C—26H₃, 27H₃), 1.23 (broad singlet, C—21H₃), 0.6 (s, C—18H₃); IR ν_{max}

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3500 (s), 3300 (s), 2950 (s), 2900 (sh), 1620 (w), 1380 (s), 1220 (s), 1070 (s), 1050 (s), 1030 (s), 1000 (s), cm^{-1} ; mass spec. molecular ion m/e = 450; (analysis found: % C, 69.05; H, 9.44; S, 7.13; $\text{C}_{26}\text{H}_{42}\text{O}_4\text{S}$ requires: % C, 69.29; H, 9.39; S, 7.12).

Example 28

23-oxa-9,10-seco- 3β ,25-dihydroxy-cholesta-5(E),7(E),10(19)-triene

The silyl ether from Example 51 (160 mg) was stirred with $n\text{-Bu}_4\text{NF}$ (1 M soln. in THF, 1 ml) in refluxing THF (5 ml) for 40 min. Dilution with CH_2Cl_2 , followed by aqueous work-up and purification by plc gave the title compound (102 mg, 82%). UV λ_{max} 274 nm; $^1\text{Hnmr}$ δ 6.47 and 5.85 (ABq, $J=11\text{Hz}$, C—6H, 7H), 4.9 (s, C—19H), 4.63 (s, C—19H), 3.83 (m, W=18Hz, C—3H), 3.58—3.07 (m, C—22H₂), 3.18 (s, C—24H₂), 1.2 (s, C—26H₃, 27H₃), 1.03 (d, J=6Hz, C—21H₃), 0.58 (s, C—18H₃).

Example 29

The 3-(3'-5'-dinitrobenzoate) ester of the product of Example 28 was prepared as described previously for Ex. 6(2). Crystalline from ether/hexane. m.p. 75—77°C; $[\alpha]_D = +176^\circ$ ($c = 0.58$); $^1\text{Hnmr}$ δ 9.15 (m, 3H, aryl), 6.58 and 5.78 (ABq, $J=11\text{Hz}$, C—6H, 7H), 5.3 (m, W=12Hz, C—3H), 5.03 (s, C—19H), 4.73 (s, C—19H), 3.57—3.07 (m, C—22H₂), 3.2 (s, C—24H₂), 1.22 (s, C—26H₃, 27H₃), 1.02 (d, J=6Hz, C—21H₃), 0.47 (s, C—18H₃); IR ν_{max} 3500 (m), 2950 (s), 2900 (sh), 1730 (s), 1640 (m), 1550 (s), 1460 (m), 1340 (s), 1270 (s), 1165 (m), cm^{-1} ; mass spec. molecular ion m/e = 596; (analysis found: % C, 66.31; H, 7.55; N, 4.56; $\text{C}_{33}\text{H}_{44}\text{O}_8\text{N}_2$ requires: % C, 66.42; H, 7.43; N, 4.70).

Example 30

23-oxa-9,10-seco- 3β -(t-butyldimethylsilyloxy)-25-hydroxy-cholesta-5(Z),7(E),10(19)-triene

The corresponding 5(E) compound from Example 51 (160 mg) in benzene (30 ml) and triethylamine (3 drops) containing phenazine (35 mg) was thoroughly degassed and irradiated as described above for 30 min. Purification by plc gave (273a) (138 mg, 86%). UV max = 263 nm; $^1\text{Hnmr}$ δ 6.25 and 6.0 (ABq, $J=11\text{Hz}$, C—6H, 7H), 5.05 (s, C—19H), 4.82 (s, C—19H), 3.92 (m, W=18Hz, C—3H), 3.62—3.10 (m, C—22H₂), 3.20 (s, C—24H₂), 1.23 (s, C—26H₃, 27H₃), 1.03 (d, J=6Hz, C—21H₃), 0.91 (s, t-Bu), 0.58 (s, C—18H₃), 0.05 [s, (Si—CH₃)₂].

Example 31

23-oxa-9,10-seco- 3β ,25-dihydroxy-cholesta-5(Z),7(E),10(19)-triene

The corresponding 3-t-butyldimethylsilyl ether from Example 30 (138 mg) was stirred with $n\text{-Bu}_4\text{NF}$ (1 M soln. in THF, 2 ml) in refluxing THF (5 ml). After 45 min, the mixture was diluted with CH_2Cl_2 . Aqueous work-up followed by purification by plc gave the diol (273b) (91 mg, 85%) as an oil. UV λ_{max} 263 nm; $^1\text{Hnmr}$ δ 6.24 and 6.04 (ABq, $J=11\text{Hz}$, C—6H, 7H), 5.03 (s, C—19H), 4.83 (s, C—19H), 3.92 (m, W=18Hz, C—3H), 3.57—3.12 (m, C—22H₂), 3.25 (s, C—24H₂), 1.22 (s, C—26H₃, 27H₃), 1.03 (d, J=6Hz, C—21H₃), 0.57 (s, C—18H₃).

Example 32

The 3-(3',5'-dinitrobenzoate) ester of the product of Example 31 was prepared as in Example 6(2). Crystalline from ether/hexane m.p. 136—138°C; $[\alpha]_D = +101^\circ$ ($c = 0.615$); $^1\text{Hnmr}$ δ 9.12 (m, 3H, aryl), 6.22 and 6.01 (ABq, $J = 11\text{Hz}$, C—6H, 7H), 5.23 (m, W=18Hz, C—3H), 5.1 (s, C—19H), 4.92 (s, C—19H), 3.57—3.1 (m, C—22H₂), 3.2 (s, C—24H₂), 1.22 (s, C—26H₃, 27H₃), 1.05 (d, J=6Hz, C—21H₃), 0.53 (s, C—18H₃); IR ν_{max} 3550 (m), 2950 (s), 2900 (sh), 1750 (s), 1650 (m), 1555 (s), 1470 (m), 1350 (s), 1280 (s), cm^{-1} ; mass spec. molecular ion m/e = 596; (analysis found: % C, 66.34; H, 7.37; N, 4.61; $\text{C}_{33}\text{H}_{44}\text{O}_6\text{N}_2$ requires: % C, 66.42; H, 7.43; N, 4.70).

Example 33

23-oxa-9,10-seco- 3β -(t-butyldimethylsilyloxy)-25-(triethylsilyloxy)-cholesta-5(E),7(E),10(19)-triene

The 25-hydroxy compound from Example 51 (300 mg) in CH_2Cl_2 (10 ml) was treated with triethylsilylchloride (130 μl) in the presence of imidazole (200 mg) for 16 hrs. Acid work-up gave the title bis silylated calciferol (274) which was used in the next step without further purification.

Example 34

23-oxa-9,10-seco-3 α -hydroxy- 3β -(t-butyldimethylsilyloxy)-25-(triethylsilyloxy)-cholesta-5(E),7(E),10(19)-triene

Selenium dioxide (60 mg) was stirred in methanol (4 ml) for 45 mins. NMO (300 mg) was stirred in CH_2Cl_2 (4 ml) in the presence of anhydrous MgSO_4 for 30 min. The NMO solution was filtered into a solution of the bis silyl ether from Example 33 in 1,2-dichloroethane (4 ml) and the mixture warmed to reflux. To this refluxing mixture was added the SeO_2 /methanol mixture. After 23 min, the heating mantle was removed and the product worked up and isolated as described previously to give 190 mg [51% based on Ex. 52] of the title 1 α -hydroxylated product. UV λ_{max} 274 nm; $^1\text{Hnmr}$ δ 6.55 and 5.88 (ABq, $J=12\text{Hz}$, C—6H, 7H), 5.1 (s, C—19H), 5.0 (s, C—19H), 4.75—4.02 (m, C—1H, 3H), 3.65—3.12 (m, C—22H₂), 3.25 (s, C—24H₂).

Example 35**23-oxa-9,10-seco-1 α ,3 β ,25-trihydroxy-cholesta-5(E),7(E),10(19)-triene**

The bis silyl ether from Example 34 (190 mg) in THF (6 ml) was refluxed with nBu₄NF (1 M solution in THF, 2 ml) for 1 hr. The mixture was diluted with CH₂Cl₂. Aqueous work-up gave the title triol (103 mg, 84%) after purification by plc. Crystalline from chloroform/hexane. m.p. 141–144°C, [α]_D = +144° (c = 0.355); UV λ_{max} 272 nm (20554); ¹Hnmr (400 MHz) δ 6.58 (d, J=12Hz), 5.89 (d, J=12Hz), 5.13 (s, C—19H), 4.98 (s, C—19H), 4.50 (m, W=12Hz, C—1H), 4.26 (m, W=20 Hz, C—3H), 3.43 (m, 1H), 3.30–3.15 (m, C—22H₂, 24H₂), 1.20 (s, C—26H₃, 27H₃), 1.02 (d, J=6Hz, C—21H₃), 0.58 (s, C—18H₃); IR v_{max} 3500 (s), 2950 (s), 2900 (sh), 1640 (w), 1450 (m), 1380 (m), 1360 (m), 1045 (s), cm⁻¹; mass spec. molecular ion m/e = 418; (analysis found: % C, 74.76; H, 10.33; C₂₆H₄₂O₄ requires: % C, 74.60; H, 10.11).

Example 36**23-oxa-9,10-seco-1 α -hydroxy-3 β -(t-butyldimethylsilyloxy)-25-(triethylsilyloxy)-cholesta-5(Z),7(E),10(19)-triene**

The corresponding 5(E) compound from Example 35 (200 mg) in benzene (35 ml) containing phenazine (40 mg) and triethylamine (4 drops) was irradiated with the Hanovia lamp as described previously for 35 min to give, after purification by plc, 155 mg (78%) of the title compound as a less polar, oily product. UV λ_{max} 263 nm; ¹Hnmr δ 6.30 and 6.01 (ABq, J=12Hz, C—6H, 7H), 5.23 (s, C—19H), 4.97 (s, C—19H), 4.67–3.9 (m, C—1H, 3H), 3.53–3.07 (m, C—22H₂), 3.17 (s, C—24H₂).

Example 37**23-oxa-9,10-seco-1 α ,3 β ,25-trihydroxy-cholesta-5(Z),7(E),10(19)-triene**

The bis silyl ether from Example 36 (155 mg) and n-Bu₄NF (1 M soln. in THF, 2 ml) were stirred together in refluxing THF (5 ml) for 1 hr. Dilution with CH₂Cl₂ followed by aqueous work-up and purification by plc gave the title triol (252a) (77 mg, 77%). Crystalline from ether/hexane. m.p. 121–123°C; [α]_D = +47° (c = 0.6); UV λ_{max} 264 nm (17200); ¹Hnmr δ 6.37 and 6.05 (ABq, J=11Hz, C—6H, 7H), 5.33 (s, C—19H), 5.0 (s, C—19H), 4.57–3.87 (m, C—1H, 3H), 3.6–3.1 (m, C—22H₂), 3.23 (s, C—24H₂), 1.23 (s, C—26H₃, 27H₃), 1.05 (d, J=6Hz, C—21H₃), 0.58 (s, C—18H₃); IR v_{max} 3500 (s), 2950 (s), 2900 (sh), 1640 (w), 1450 (m), 1380 (m), 1360 (m), 1045 (s), cm⁻¹; mass spec. molecular ion m/e = 418; (analysis found: % C, 74.47; H, 9.97; C₂₆H₄₂O₄ requires: % C, 74.60; H, 10.11).

Example 38**9,10-seco-3 β -(triethylsilyloxy)-20(S)-(p-toluenesulphonyloxymethyl)-pregna-5(E),7(E),10(19)-triene****Method A**

To the hydroxy compound from Example 13(D) (400 mg) in pyridine (5 ml) was added tosylichloride (350 mg) and the mixture stirred overnight at room temperature. Water was added and the mixture diluted with ether. Acid work-up gave, after purification by plc, 310 mg (58%) of the title tosylate.

¹Hnmr δ 7.73 (d, J=8Hz, 2H, aryl), 7.28 (d, J=8Hz, 2H, aryl), 6.43 and 5.81 (ABq, J=11Hz, C—6H, 7H), 4.92 (s, C—19H), 4.63 (s, C—19H), 4.2–3.57 (m, C—3H, 22H₂), 2.48 (s, aryl-CH₃); IR v_{max} (thin film) 2960 (s), 2900 (sh), 1600 (w), 1460 (m), 1360 (s), 1190 (s), 1175 (s), 1090 (s), cm⁻¹.

Method B

The crude SO₂ adducts of 9,10-seco-3 β -triethylsilyloxy-20(S)-(hydroxymethyl)-pregna-5(E),7(E),10(19)-triene from Example 12(1) (3.2 g) was stirred overnight in pyridine (40 ml) at 5°C with p-toluenesulphonyl chloride (4 g). The mixture was cooled to 0°C, water added and, after a few minutes, the mixture was diluted with Et₂O. After an acid work-up, the crude oily product (281) was taken up in ethanol (100 ml) and refluxed in the presence of NaHCO₃ (4 g) for 1 hr. The mixture was concentrated and partitioned between CH₂Cl₂/water/brine. The organic solution was dried and chromatographed to give 2.64 g (70%) of the required vitamin (278c) nmr and IR identical to the product obtained by Method A.

Example 39**9,10-seco-3 β -hydroxy-20(S)-[fluoromethyl]-pregna-5(E),7(E),10(19)-triene**

The tosylate from Example 38 (200 mg) in THF (5 ml) was refluxed for 45 min in the presence of n-Bu₄NF (1 M soln. in THF, 1 ml). The mixture was diluted with CH₂Cl₂. Aqueous work-up followed by purification by plc gave 70 mg (63%) of the title fluoride (279). ¹Hnmr δ 6.5 and 5.83 (ABq, J=11Hz, C—6H, 7H), 4.97 (s, C—19H), 4.7 (br, s, C—19H, 22H), 4.2–3.6 (m, C—3H, 22H), 1.1 (d, J=6Hz, C 21H₃), 0.6 (s, C—18H₃).

Example 40**9,10-seco-1 α -hydroxy-3 β -(triethylsilyloxy)-20(S)-(p-toluenesulphonyloxymethyl)-pregna-5(E),7(E),10(19)-triene**

Selenium dioxide (56 mg) was stirred in acetonitrile (3.5 ml) for 45 min. NMO (280 mg) was stirred in CH₂Cl₂ (3.5 ml) in the presence of anhydrous MgSO₄ for 30 min. The NMO solution was filtered into a solution of the 1-desoxy compound from Example 387 (308 mg) in 1,2-dichloroethane (3.5 ml) and the mixture warmed to reflux. To this was added the SeO₂/CH₂CN mixture, and refluxing continued for a further

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5.5 min. The reaction mixture was cooled in an ice bath, diluted with CH_2Cl_2 and worked up as previously to give 180 mg (57%) of the title 1-hydroxy compound. $^1\text{Hnmr}$ δ 7.73 (d, $J=8\text{Hz}$, 2H, aryl), 7.28 (d, $J=8\text{Hz}$, 2H, aryl), 6.43 and 5.81 (ABq, $J=11\text{Hz}$, C—6H, 7H), 5.03 (s, C—19H), 4.93 (s, C—19H), 4.63—3.6 (m, C—1H, 3H, 22H₂), 2.48 (s, aryl-CH₃).

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Example 41

9,10-seco-1 α ,3 β -dihydroxy-20(S)-(p-toluenesulphonyloxy)methyl)-pregna-5(E),7(E),10(19)-triene

The 3-triethylsilylether derivative from Example 40 (180 mg) in THF (5 ml) containing n-Bu₄NF (1 M soln. in THF, 0.4 ml) was stirred for 15 min. The mixture was diluted with CH_2Cl_2 . An aqueous work-up and purification by plc gave 118 mg (81%) of the title diol. Solid from CH_2Cl_2 /hexane. m.p. 97—99°C; $[\alpha]_D = +132^\circ$ ($c = 0.57$); UV λ_{max} 272 nm (23360) and 218 nm (15920); $^1\text{Hnmr}$ δ 7.73 (d, $J=8\text{Hz}$, 2H, aryl), 7.28 (d, $J=8\text{Hz}$, 2H, aryl), 6.43 and 5.81 (ABq, $J=11\text{Hz}$, C—6H, 7H), 5.03 (s, C—19H), 4.93 (s, C—19H), 4.63—3.53 (m, C—1H, 3H, 22H₂), 2.5 (s, aryl-CH₃), 1.02 (d, $J=6\text{Hz}$, C—21H₃), 0.57 (s, C—18H₃); IR ν_{max} 3500 (s), 2950 (s), 2900 (sh), 1600 (w), 1450 (m), 1355 (s), 1190 (s), 1175 (s), cm^{-1} .

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Example 42

9,10-seco-1 α -hydroxy-3 β -(triethylsilyloxy)-20(S)-(p-toluenesulphonyloxy)methyl)-pregna-5(Z),7(E),10(19)-triene

The corresponding 5(E) compound from Example 40 (225 mg) in benzene (35 ml) containing triethylamine (3 drops) was irradiated as described above with anthracene (45 mg) as triplet sensitizer for 30 min to give, after plc, 185 mg (82%) of the title compound. UV λ_{max} 263 nm and 216 nm; $^1\text{Hnmr}$ δ 7.73 (d, $J=8\text{Hz}$, 2H, aryl), 7.3 (d, $J=8\text{Hz}$, 2H, aryl), 6.28 and 5.98 (ABq, $J=11\text{Hz}$, C—6H, 7H), 5.28 (s, C—19H), 4.92 (s, C—19H), 4.55—3.58 (m, C—1H, 3H, 22H₂), 2.45 (s, aryl-CH₃).

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Example 43

9,10-seco-1 α ,3 β -dihydroxy-20(S)-(p-toluenesulphonyloxy)methyl)-pregna-5(Z),7(E),10(19)-triene

The silyl ether from Example 43 (185 mg) in THF (5 ml) containing n-Bu₄NF (1 M soln. in THF, 0.32 ml) was stirred for 15 min at room temperature. Dilution with CH_2Cl_2 aqueous work-up and purification by plc gave the title diol (110 mg, 73%). UV λ_{max} 263 nm (17427) and 216 nm (18672); $^1\text{Hnmr}$ δ 7.68 (d, $J=8\text{Hz}$, 2H, aryl), 7.23 (d, $J=8\text{Hz}$, 2H, aryl), 6.28 and 5.97 (ABq, $J=11\text{Hz}$, C—6H, 7H), 5.27 (s, C—19H), 4.93 (s, C—19H), 4.57—3.6 (m, C—1H, 3H, 22H₂), 2.45 (s, aryl-CH₃), 1.05 (d, $J=6\text{Hz}$, C—21H₃), 0.52 (s, C—18H₃).

Example 44

1-amino-2-methyl-2-hydroxy-propane

To a well-stirred mixture of lithium aluminium hydride (12 g) in ether (200 ml) at 0°C was added dropwise over 1 hr a solution of acetone cyanohydrin (11.2 g, 12 ml) in ether (50 ml). The mixture was stirred at room temperature overnight. After cooling to 0°C, water (24 ml) was cautiously added dropwise. After the quenching was complete, anhydrous Na₂SO₄ (65 g) was added and stirring at room temperature was continued for a further 2.5 hr. The solid was filtered off and the ether evaporated to give, after distillation, 4.8 g (41%) of the title compound as a viscous, colourless liquid.
b.p. 74—76°C/14 mm Hg (lit.¹⁶⁹ 62—64°C/13 mm Hg) $n_D^{20} = 1.4463$ (lit.¹⁶⁹ $n_D^{20} = 1.4467$); $^1\text{Hnmr}$ δ 2.6 (s, 2.6 (s, 2H), 1.87 (s, 3H, exchanges with D₂O), 1.2 (s, 6H); IR ν_{max} (thin film) 3400 (s), 3000 (m), 1600 (m), 1475 (m), 1380 (m), 1360 (m), 1220 (m), 1170 (m), 1110 (m), 960 (m), cm^{-1} .

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Example 45

23-aza-9,10-seco-1 α ,3 β ,25-trihydroxy-cholesta-5(Z),7(E),10(19)-triene

A solution of the tosylate from Example 44 (100 mg) in 1-amino-2-methyl-2-hydroxy-propane (0.5 ml) was degassed and then stirred under argon at 50—55°C for 6 hr and then at room temperature for a further 12 hr. The solution was diluted with CH_2Cl_2 and washed with water/brine and dried to give, after purification by plc, 44 mg (53%) of the title triol. $[\alpha]_D = +24^\circ$; UV λ_{max} 264 nm (15400); $^1\text{Hnmr}$ δ 6.38 and 6.07 (ABq, $J=11\text{Hz}$, C—6H, 7H), 5.35 (s, C—19H), 5.02 (s, C—19H), 4.67—3.93 (m, C—1H, 3H), 2.5 (s, C—24H₂), 1.2 (s, C—26H₃, 27H₃), 1.02 (d, $J=6\text{Hz}$, C—21H₃), 0.57 (s, C—18H₃); IR ν_{max} 3500 (s), 2950 (s), 2900 (sh), 1640 (w), 1460 (m), 1380 (m), 1055 (m), cm^{-1} ; mass measurement found: 417.3242; C₂₈H₄₃O₃N requires: 417.3243.

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Example 46

23-aza-9,10-seco-1 α ,3 β ,25-trihydroxy-cholesta-5(Z),7(E),10(19)-triene-23-N-acetyl

The crude amine from Example 45 derived from the tosylate (100 mg) as described above, in methanol (5 ml) containing K₂CO₃ (500 mg) was treated with acetic anhydride (0.2 ml). The mixture was diluted with CH_2Cl_2 washed with brine and dried to give, after plc, 50 mg [55% fr m tosylate] of the title amide. Solid from CH_2Cl_2 /hexane. m.p. 107—109°C; $[\alpha]_D = -14^\circ$ ($c = 0.49$); UV λ_{max} 263 nm (16275); $^1\text{Hnmr}$ δ 6.37 and 6.05 (ABq, $J=11\text{Hz}$, C—6H, 7H), 5.33 (s, C—19H), 5.0 (s, C—19H), 4.65—4.02 (m, C—1H, 3H), 3.4 (s, C—24H₂), 2.17 (s, acetyl), 1.22 (s, C—26H₃, 27H₃), 0.95 (d, $J=7\text{Hz}$, C—21H₃), 0.6 (s, C—18H₃); IR ν_{max} 3550 (s), 2950 (s), 2900 (sh), 1640 (s), 1460 (m), 1380 (m), 1055 (m), cm^{-1} ; (analysis found: % C, 70.80; H, 10.12; N, 2.77; C₂₈H₄₅O₄N requires: C, 73.16; H, 9.87; N, 3.05; C₂₈H₄₅O₄N · H₂O requires: % C, 70.40; H, 9.92; N, 2.93).

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Example 47

9,10-s co- β ,25-dihydroxy-cholesta-5(E),7(E),10(19)-triene

Magnesium turning were washed with diluted HCl/water/acetone/ether and dried in vacuo for 24 hr. The 1-bromo-4-methyl-4-triethylsilylbutane (1 g) in freshly distilled (from LiAlH₄) THF (10 ml) containing magnesium metal (82 mg) was refluxed for 2 hr.

Cuprous iodide (100 mg) was placed in a flask and purged with argon, whilst cooling to 0°C. To this was added the above Grignard solution (5 ml), and the purple coloured mixture stirred for an additional 30 min at 0°C. A solution of the tosylate (278c) (200 mg) in ether (2 ml) was added and the mixture stirred for 40 min at room temperature. Water was added and the reaction mixture extracted with ether. After an acid work-up, the non-polar product was isolated by plc contaminated with large quantities of low molecular weight alkyl residues. This mixture was stirred with n-Bu₄NF (1 M soln. in THF, 2 ml) in refluxing THF (5 ml) for 2 hr. Dilution with CH₂Cl₂ followed by aqueous work-up and purification by plc gave 110 mg [82% from tosylate (278c)] of this previously described title diol. The physical and spectral properties of this material were identical in all respects to the product obtained from the phthalazine adduct.

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Example 48

9,10-seco- β ,25-dihydroxy-cholest-5(Z),7(E),10(19)-triene

The product from Example 47 (100 mg) in benzene (30 ml) and triethylamine (3 drops) containing anthracene (25 mg) was thoroughly degassed and irradiated for 1 hr as described above to give, after purification by plc, the title 5(Z) compound (90 mg, 82%). The physical and spectral properties of this material were identical in all respects to the product obtained via the phthalazine adduct. A mixed melting point determination of this material and an authentic sample, kindly supplied by Roussel Uclaf, Inc. (Romainville, France) was undepressed.

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Example 49

9,10-seco-1 α ,3 β -bis(triethylsilyloxy)-20(S)-(p-toluenesulphonyloxymethyl)-pregna-5(Z),7(E),10(19)-triene

The tosylate (276b) (105 mg) in CH₂Cl₂ (5 ml) containing imidazole (75 mg) and triethylsilylchloride (45 μ l) was stirred at room temperature for about 15 min. Water was added and the mixture diluted with CH₂Cl₂. Acid work-up gave the non-polar title bis silyl ether which was used without further purification.

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Example 50

9,10-seco-1 α ,3 β ,25-trihydroxy-cholesta-5(Z),7(E),10(19)-triene

To the alkyl copper reagent at 0°C prepared exactly as described above, was added a solution of the above tosylate (276c) in THF (3 ml) and the mixture stirred at room temperature for 25 min. Work-up and purification as in Example 6(1) gave the tris triethylsilyl derivative contaminated with large quantities of low molecular weight alkyl residues. This mixture was treated with n-Bu₄NF (1 M soln. in THF, 4 ml) in THF (5 ml) for 20 min at room temperature followed by 1.5 hr at reflux to give, after the usual work-up and purification by plc, a mixture of the title steroidial triol [(38 mg, 63% from (276b))] contaminated with isopentane diol (10 mg). Dissolution of this mixture in CHCl₃ gave the required product as its crystalline CHCl₃ solvate (25 mg). m.p. 99–105°C (lit. 106–112°C¹⁴², 103–106°C¹³⁸); [α]_D (Et₂O) = +35° (c = 0.86); UV λ_{max} 264 nm (16820); ¹Hnmr δ (acetone-d₆) 8.07 (s, CHCl₃), 6.35 and 6.18 (ABq, J=12Hz, C—6H, 7H), 5.38 (s, C—19H), 4.93 (s, C—19H), 4.7–4.07 (m, C—1H, 3H), 1.2 (s, C—26H₃, 27H₃), 1.0 (broad singlet, C—21H₃), 0.6 (s, C—18H₃); IR v_{max} 3500 (s), 2950 (s), 2900 (sh), 1640 (w), 1480 (m), 1440 (m), 1380 (m), 1360 (m), 1140 (m), 1050 (s).

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Example 51

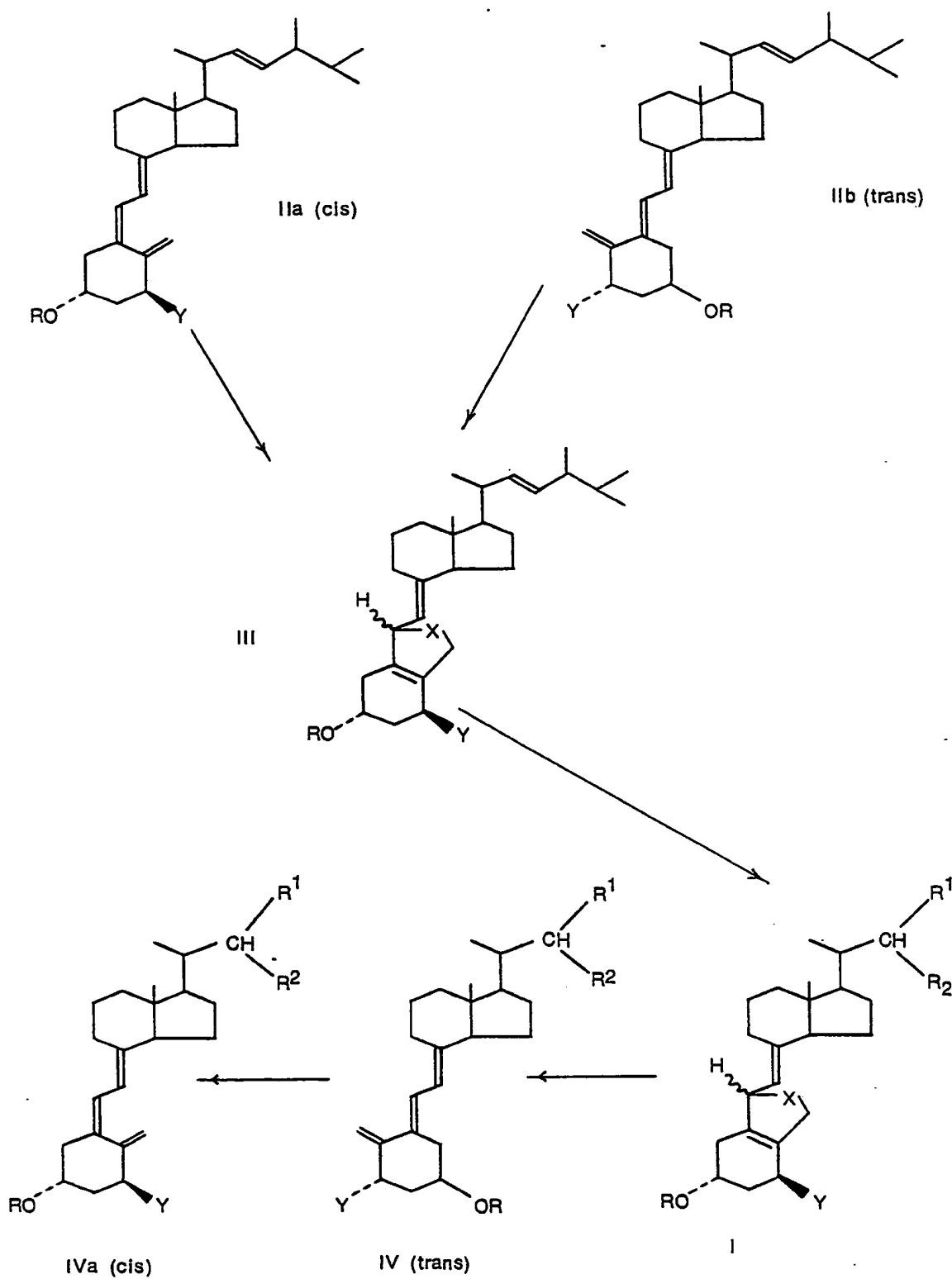
23-oxa-9,10-seco-3 β -(t-butylidimethylsilyloxy)-25-hydroxy-cholesta-5(E),7(E),10(19)-triene

The 22-hydroxy compound (167b) (425 mg) in benzene (5 ml) was refluxed with isobutylene epoxide (1 ml) in the presence of dibenzo-18-crown-6 (100 mg) and potassium t-butoxide (500 mg) for 55 min. Water was added and the mixture diluted with CH₂Cl₂. The organic layer was washed with aqueous K₃PO₄/water/5% aqueous NaHCO₃/brine and dried. Purification by plc gave 330 mg (67%) of the slightly less polar oily product. ¹Hnmr δ 6.45 and 5.85 (ABq, J=12Hz, C—6H, 7H), 4.9 (s, C—19H), 4.63 (s, C—19H), 3.92 (m, W=18Hz, C—3H), 3.63–3.12 (m, C—22H₂), 3.22 (s, C—24H₂), 1.23 (s, C—26H₃, 27H₃), 1.05 (d, J=6Hz, C—21H₃), 0.92 (s, t-Bu), 0.6 (s, C—18H₃), 0.05 s, [(Si—CH₃)₂].

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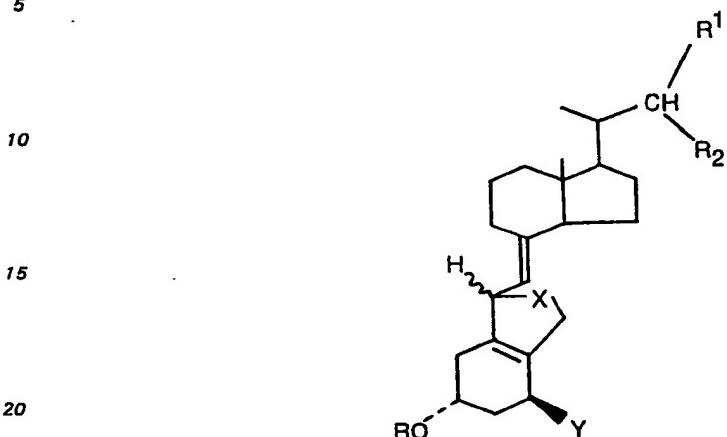
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Claims for the Contracting States: BE CH DE FR GB IT LI LU NL SE

1. Compounds of the general formula

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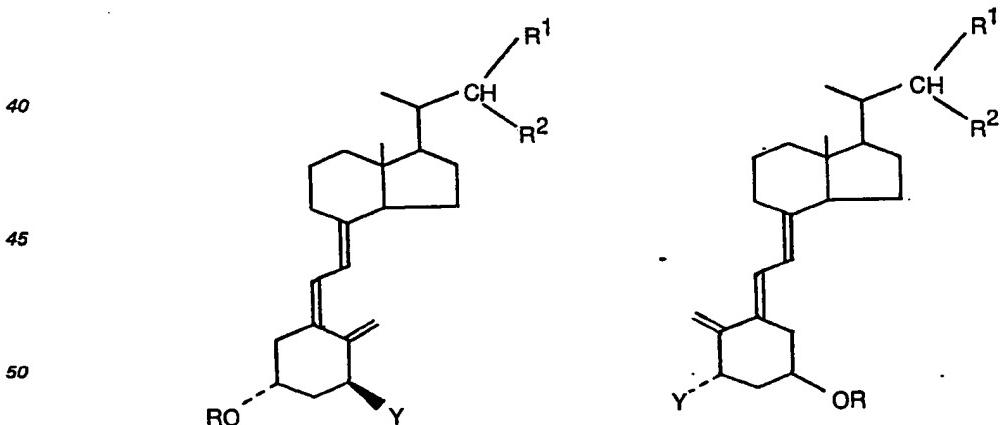


wherein R represents a hydrogen atom or a hydroxyl protecting group, Y represents a hydrogen atom of an
 25 optionally protected hydroxyl group, X represents SO_2 or the residue of a diacylazo dienophile and either
 R^1 represents a halogen atom a hydrocarbysulphonyloxy group or a group of the formula $\text{Z}-R^3$ (in
 which Z represents O , S , SO , NR^4 or CR^4R^6 and R^3 , R^4 and R^5 , which may be the same
 30 or different, each represent a hydrogen atom or a straight or branched aliphatic group having 1—12 carbon
 atoms and which may optionally carry one or more substituents) and R^2 represents a hydrogen atom or R^1
 and R^2 together represent an oxo group or an optionally substituted alkylidene group, except that R^1 and R^2
 together with the group $-\text{CH}(\text{CH}_3)\text{CH}-$ to which they are attached do not represent a group having the
 branched 17 β -hydrocarbyl side chain skeleton of vitamin D₂ or vitamin D₃.

2. Compounds as claimed in claim 1 in which the dienophile is a cyclic diacylazo compound.

3. Compounds of general formula IV or IVa,

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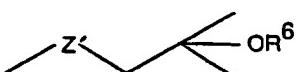
IVa (cis)

IV (trans)

wherein R, Y, R¹ and R² are as defined in claim 1.

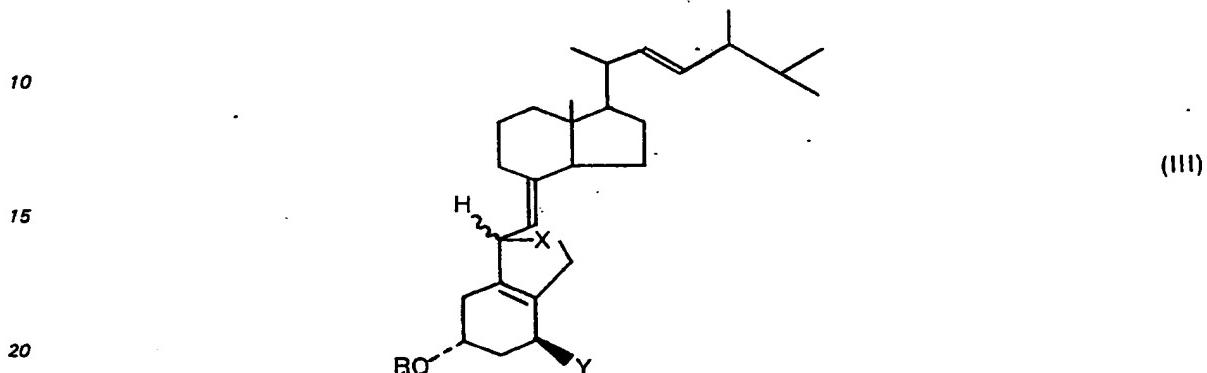
4. Compounds of general formulae I, IV or IVa as claimed in any one of claims 1 to 3 wherein R¹
 60 represents a halogen atom, a hydroxyl or tosyloxy group or a group of formula:

65



(in which Z' represents —O—, —S—, —NH— or —SO— and R⁶ represents a hydrogen atom or a hydroxyl protecting group) and R² represents a hydrogen atom or R¹ and R² together represent an alkylidene group having 1 to 8 carbon atoms optionally substituted by one or more substituents selected from halogen atoms and optionally protected hydroxyl groups.

5. A process for the preparation of compounds of general formula I as defined in claim 1 in which R¹ and R² together represent an oxo group which comprises subjecting a compound of formula III,



(wherein R, Y and X are as defined in claim 1) to oxidative cleavage.

6. A process as claimed in claim 5 wherein the aldehyde of formula I so formed is subsequently reduced to give a compound of formula I wherein R¹ represents a hydroxyl group.
7. A process as claimed in claim 5 wherein the aldehyde of formula I so formed is subsequently reacted with a Wittig reagent to give a compound of formula I wherein R¹ and R² together represent an optionally substituted alkylidene group, the double bond of which may then, if desired, be reduced.

8. A process for the preparation of compounds of general formula IV or IVa as defined in claim 3 which comprises deprotecting a corresponding compound of formula I as defined in claim 1 by removal of the residue X and subsequently, optionally after conversion of the group R¹ to another group R¹, isomerising the compound of formula IV thus obtained to a compound of formula IVa.

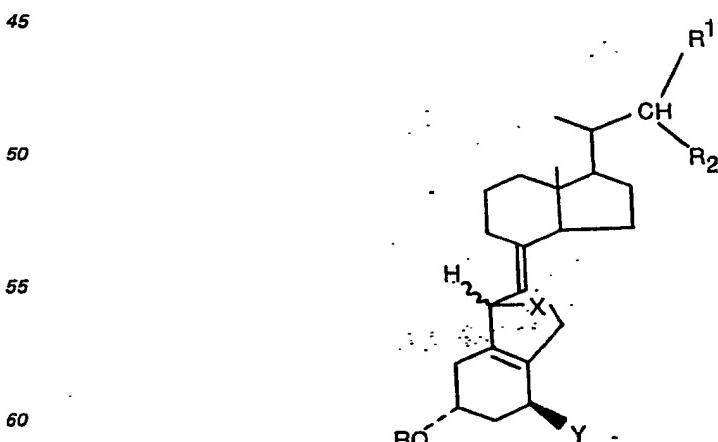
9. A process as claimed in claim 6 or claim 8 wherein a product is obtained in which R¹ represents a hydroxyl group and the said hydroxyl group is converted into a halogen atom or a hydrocarbylsulphonyloxy group.

10. A process as claimed in any one of claims 6, 8 and 9 wherein a product is obtained in which R¹ represents a halogen atom or a hydroxyl or hydrocarbylsulphonyloxy group which product is converted into a product wherein R¹ represents a group of formula —ZR³ as defined in claim 1 other than a hydroxyl group.

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Claims for the Contracting State: AT

1. A process for the preparation of compounds of the general formula



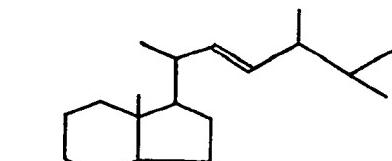
(wherein R represents a hydrogen atom or a hydroxyl protecting group, Y represents a hydrogen atom or an optionally protected hydroxyl group, X represents —SO₂ or the residue of a diacylazodienophile and either R¹ represents a halogen atom, a hydrocarbylsulphonyloxy group or a group of the formula —Z—R³

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(in which Z represents $-O-$, $-S-$, $-SO-$, $-NR^4-$ or $-CR^4R^5-$ and R^3 , R^4 and R^5 , which may be the same or different, each r present a hydrogen atom or a straight or branched aliphatic group having 1-12 carbon atoms and which may optionally carry one or more substituents) and R^2 represents a hydrogen atom or R^1 and R^2 together represent an oxo group or an optionally substituted alkylidene group, except

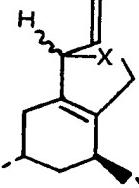
- 5 that R^1 and R^2 together with the group $-CH(CH_3)CH-$ to which they are attached do not represent a group having the branched 17β -hydrocarbyl side chain skeleton of vitamin D_2 or vitamin D_3) which comprises subjecting a compound of formula III,

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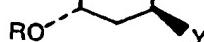


(III)

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(wherein R, Y and X are as defined above) to oxidative cleavage to form an aldehyde of formula I (in which R^1 and R^2 together represent an oxo group) and subsequently, if desired

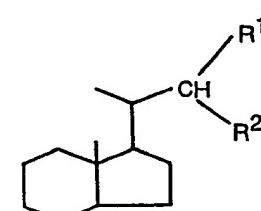
- 30 either reacting the said aldehyde of formula I with a Wittig reagent to give a compound of formula I wherein R^1 and R^2 together represent an optionally substituted alkylidene group, the double bond of which may then, if desired, be reduced.

or reducing the said aldehyde of formula I to give a compound of formula I wherein R^1 represents a hydroxyl group, the said hydroxyl group being then optionally converted into a halogen atom, a hydrocarbysulphonyloxy group or a group of formula $-ZR^3$ as defined above other than a hydroxyl group.

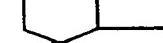
- 35 2. A process as claimed in claim 1 in which the dienophile is a cyclic diacylazo compound.

3. A process for the preparation of compounds of general formula IV or IVa,

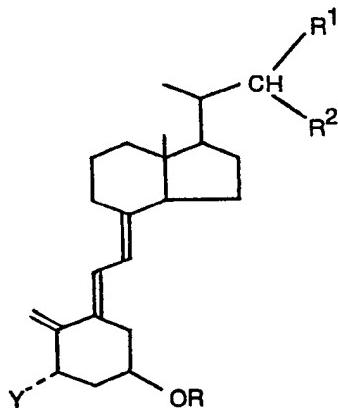
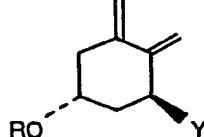
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IVa (cis)

IV (trans)

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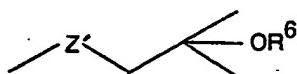
(wherein R, Y, R^1 and R^2 are as defined in claim 1) which comprises deprotecting a corresponding compound of formula I as defined in claim 1 by removal of the residue X and subsequently, optionally after conversion of the group R^1 to another group R^1' , isomerising the compound of formula IV thus obtained to a compound of formula IVa.

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4. A process as claimed in any preceding claim for the preparation of compounds of general formula I, IV or IVa wherein R¹ represents a halogen atom, a hydroxyl or tosyloxy group or a group of formula:

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(in which Z' represents —O—, —S—, —NH— or —SO— and R⁶ represents a hydrogen atom or a hydroxyl protecting group) and R² represents a hydrogen atom or R¹ and R² together represent an alkylidene group having 1 to 8 carbon atoms optionally substituted by one or more substituents selected from halogen atoms and optionally protected hydroxyl groups.

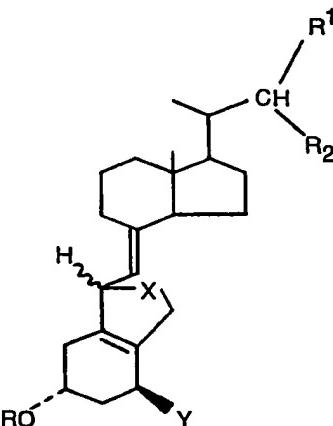
10 5. A process as claimed in any preceding claim wherein a product is obtained in which R¹ represents a hydroxyl group and the said hydroxyl group is converted into a halogen atom or a hydrocarbysulphonyloxy group.

15 6. A process as claimed in any preceding claim wherein a product is obtained in which R¹ represents a halogen atom or a hydroxyl or hydrocarbysulphonyloxy group which product is converted into a product wherein R¹ represents a group of formula —ZR³ as defined in claim 1 other than a hydroxyl group.

20 **Patentansprüche für die Vertragsstaaten: BE CH DE FR GB IT LI LU NL SE**

1. Verbindungen der allgemeinen Formel

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45 worin R ein Wasserstoffatom oder eine Hydroxyl-Schutzgruppe bedeutet, Y ein Wasserstoffatom oder eine gegebenenfalls geschützte Hydroxylgruppe darstellt, X für —SO₂ oder den Rest eines Diacylazo-Dienophils steht und entweder R¹ bedeutet ein Halogenatom oder eine Hydrocarbysulfonyloxygruppe oder eine Gruppe der Formel —Z—R³ (worin Z für —O—, —S—, —SO—, —NR⁴— oder —CR⁴R⁵— steht und R³, R⁴ und R⁵, welche gleich oder verschieden sein können, jeweils ein Wasserstoffatom oder eine gerade oder verzweigte, aliphatische Gruppe mit 1 bis 12 Kohlenstoffatomen bedeuten und die gegebenenfalls einen oder mehrere Substituenten tragen kann) und R² ein Wasserstoffatom bedeutet oder R¹ und R² bedeuten zusammen eine Oxogruppe oder eine gegebenenfalls substituierte Alkylidengruppe, mit der Ausnahme, daß R¹ und R² zusammen mit der Gruppe —CH(CH₃)CH—, an die sie gebunden sind, nicht eine Gruppe bedeuten, welche das verzweigte 17β-Hydrocarbys-Seitenkettengerüst von Vitamin D₂ oder Vitamin D₃ hat.

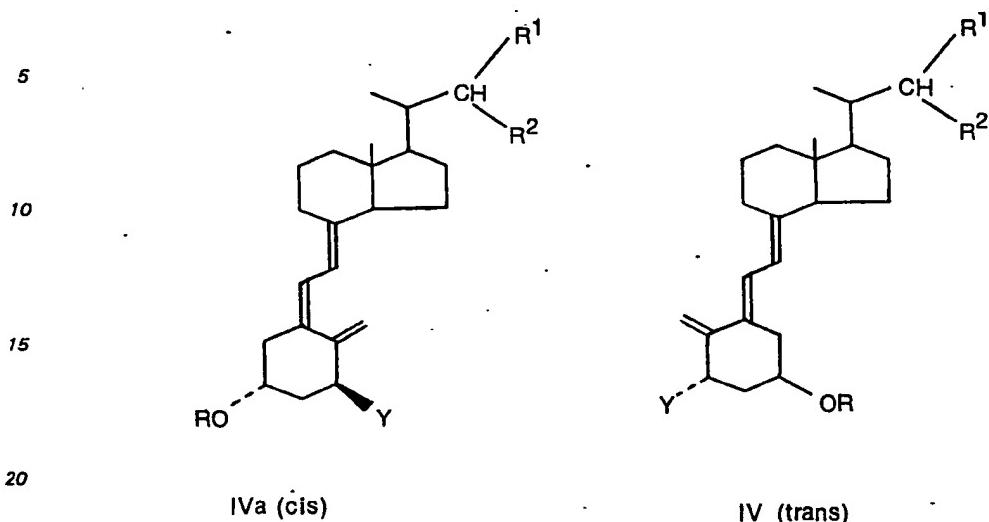
50 2. Verbindungen gemäß Anspruch 1, worin das Dienophil eine cyclische Diacylazo-Verbindung ist.

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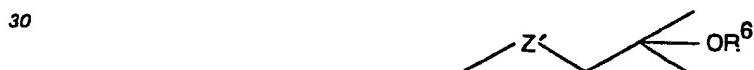
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3. Verbindungen der allgemeinen Formel IV oder IVa

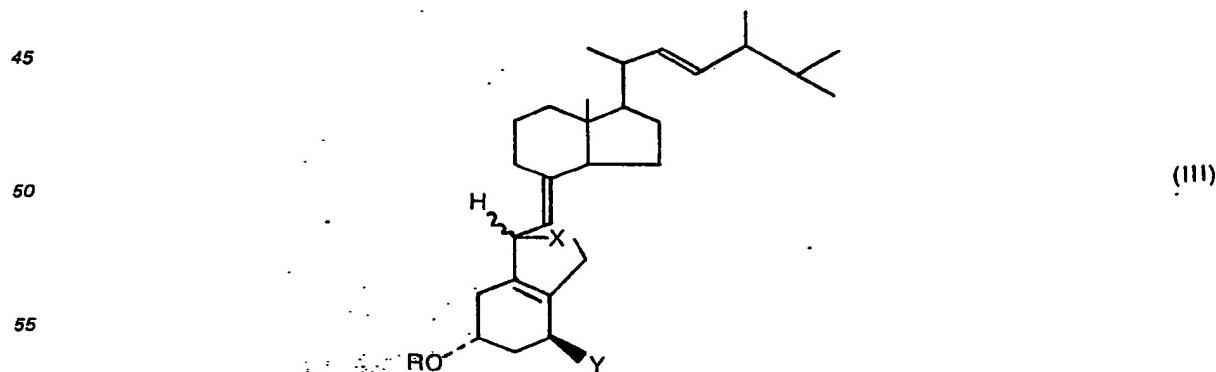


- 25 worin R, Y, R¹ und R² wie in Anspruch 1 definiert sind.
 4. Verbindungen der allgemeinen Formeln I, IV oder IVa, wie in einem der Ansprüche 1 bis 3 beansprucht, worin R¹ ein Halogenatatom, eine Hydroxyl- oder Tosyloxygruppe oder eine Gruppe der Formel



- 35 bedeutet (worin Z' für —O—, —S—, —NH— oder —SO— steht und R⁶ ein Wasserstoffatom oder eine Hydroxyl-Schutzgruppe bedeutet) und R² ein Wasserstoffatom bedeutet oder R¹ und R² bedeuten zusammen eine Alkylidengruppe mit 1 bis 8 Kohlenstoffatomen, gegebenenfalls substituiert durch einen oder mehrere Substituenten, ausgewählt aus Halogenatomen und gegebenenfalls geschützten Hydroxylgruppen.

40 5. Verfahren zur Herstellung von Verbindungen der allgemeinen Formel I, wie in Anspruch 1 definiert, worin R¹ und R² zusammen eine Oxogruppe bedeuten, dadurch gekennzeichnet, daß eine Verbindung der Formel III



- 60 (worin R, Y und X wie in Anspruch 1 definiert sind) der oxidativen Spaltung unterworfen wird.
 6. Verfahren gemäß Anspruch 5, dadurch gekennzeichnet, daß der so gebildete Aldehyd der Formel I anschließend reduziert wird, um eine Verbindung der Formel I zu ergeben, worin R¹ eine Hydroxylgruppe darstellt.

65 7. Verfahren gemäß Anspruch 5, dadurch gekennzeichnet, daß der so gebildete Aldehyd der Formel I anschließend mit einem Wittig-Reagens umgesetzt wird, um eine Verbindung der Formel I zu ergeben,

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worin R¹ und R² zusammen eine gegebenenfalls substituierte Alkylidengruppe darstellen, deren Doppelbindung dann gewünscht falls reduziert werden kann.

8. Verfahren zur Herstellung von Verbindungen der allgemeinen Formel IV oder IVa, wobei in Anspruch 3 definiert, dadurch gekennzeichnet, daß in einer entsprechenden Verbindung der Formel I, wobei in Anspruch 5 definiert, die Schutzgruppe durch Entfernung des Restes X entfernt wird und anschließend gegebenenfalls nach Überführung der Gruppe R¹ in eine andere Gruppe R¹ die so erhaltene Verbindung der Formel IV zu einer Verbindung der Formel IVa isomerisiert wird.

9. Verfahren gemäß Anspruch 6 oder Anspruch 8, dadurch gekennzeichnet, daß ein Produkt erhalten wird, worin R¹ eine Hydroxylgruppe bedeutet, und diese Hydroxylgruppe in ein Halogenatom oder eine 10 Hydrocarbysulfonyloxygruppe umgewandelt wird.

10. Verfahren gemäß einem der Ansprüche 6, 8 und 9, dadurch gekennzeichnet, daß ein Produkt erhalten wird, worin R¹ ein Halogenatom oder eine Hydroxyl- oder Hydrocarbysulfonyloxygruppe darstellt, welches Produkt in ein Produkt überführt wird, worin R¹ eine Gruppe der Formel —ZR³, wie in Anspruch 1 definiert, anders als eine Hydroxylgruppe, darstellt.

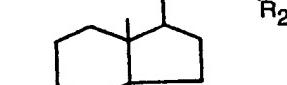
15 Patentansprüche für den Vertragsstaat: AT

1. Verfahren zur Herstellung einer Verbindung der allgemeinen Formel

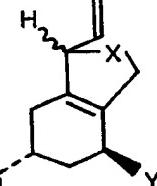
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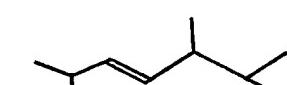


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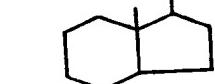
I

[worin R ein Wasserstoffatom oder eine Hydroxyl-Schutzgruppe bedeutet, Y ein Wasserstoffatom oder eine gegebenenfalls geschützte Hydroxylgruppe darstellt, X für —SO₂ oder den Rest eines Diacylazo-Dienophils steht und entweder R¹ ein Halogenatom oder eine Hydrocarbysulfonyloxygruppe oder eine Gruppe der Formel —Z—R³ bedeutet (worin Z für —O—, —S—, —SO—, —NR⁴ oder —CR⁴R⁵ steht und R³, R⁴ und R⁵, die gleich oder verschieden sein können, jeweils ein Wasserstoffatom oder eine gerade oder verzweigte, aliphatische Gruppe mit 1 bis 12 Kohlenstoffatomen bedeuten und die gegebenenfalls einen oder mehrere Substituenten tragen können) und R² ein Wasserstoffatom darstellt oder R¹ und R² bedeuten zusammen 40 eine Oxogruppe oder eine gegebenenfalls substituierte Alkylidengruppe, mit der Ausnahme, daß R¹ und R² zusammen mit der Gruppe —CH(CH₃)CH—, an die sie gebunden sind, nicht eine Gruppe bedeuten, welche 45 das verzweigte 17β-Hydrocarbyl-Seitenkettengerüst von Vitamin D₂ oder Vitamin D₃ hat], dadurch gekennzeichnet, daß eine Verbindung der Formel III

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(III)

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(worin R, Y und X wie vorstehend definiert sind) der oxidativen Spaltung zur Bildung eines Aldehyds der Formel I (worin R¹ und R² zusammen eine Oxogruppe darstellen) unterworfen wird und anschließend gewünschtenfalls

- 5 entweder dieser Aldehyd der Formel I mit einem Wittig-Reagens umgesetzt wird, um eine Verbindung der Formel I zu ergeben, worin R¹ und R² zusammen eine gegebenenfalls substituierte Alkylidengruppe darstellen, deren Doppelbindung dann gewünschtenfalls reduziert werden kann;
- 10 oder der genannte Aldehyd der Formel I reduziert wird, um eine Verbindung der Formel I zu ergeben, worin R¹ eine Hydroxylgruppe darstellt und diese Hydroxylgruppe dann gegebenenfalls in ein Halogenatom, eine Hydrocarbonylsulfonyloxygruppe oder eine Gruppe der Formel —ZR³, wie vorstehend definiert, anders als eine Hydroxylgruppe, überführt wird.

2. Verfahren gemäß Anspruch 1, dadurch gekennzeichnet, daß das Dienophil eine cyclische Diacylazo-Verbindung ist.

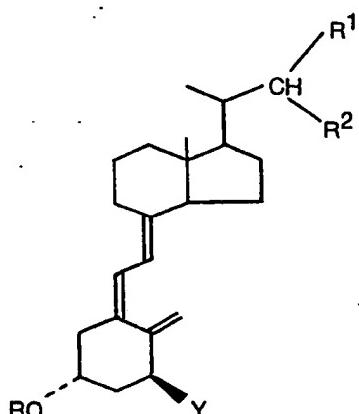
3. Verfahren zur Herstellung von Verbindungen der allgemeinen Formel IV oder IVa

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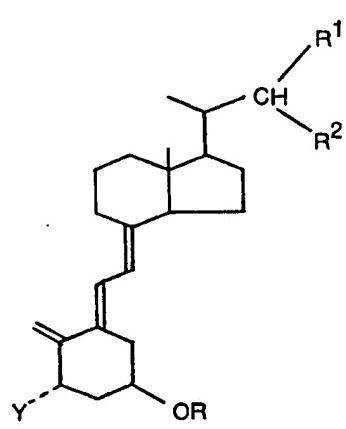
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IVa (cis)



IV (trans)

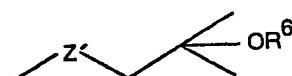
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(worin R, Y, R¹ und R² wie in Anspruch 1 definiert sind), dadurch gekennzeichnet, daß in einer entsprechenden Verbindung der Formel I, wie in Anspruch 1 definiert, die Schutzgruppe durch Entfernung des Restes X entfernt wird und anschließend, gegebenenfalls nach Umwandlung der Gruppe R¹ in eine

- 40 andere Gruppe R¹, die so erhaltene Verbindung der Formel IV zu einer Verbindung der Formel IVa isomerisiert wird.

4. Verfahren gemäß einem der vorhergehenden Ansprüche zur Herstellung von Verbindungen der allgemeinen Formeln I, IV oder IVa, worin R¹ eine Halogenatom, eine Hydroxyl- oder Tosyloxygruppe oder eine Gruppe der Formel

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bedeutet (worin Z' für —O—, —S—, —NH— oder —SO— steht und R⁶ ein Wasserstoffatom oder eine Hydroxyl-Schutzgruppe darstellt) und R² ein Wasserstoffatom bedeutet oder R¹ und R² zusammen eine Alkylidengruppe mit 1 bis 8 Kohlenstoffatomen, gegebenenfalls durch einen oder mehrere Substituenten, ausgewählt aus Halogenatomen und gegebenenfalls geschützten Hydroxylgruppen, substituiert, darstellen.

55 5. Verfahren gemäß einem der vorhergehenden Ansprüche, dadurch gekennzeichnet, daß ein Produkt erhalten wird, worin R¹ eine Hydroxylgruppe darstellt und diese Hydroxylgruppe in ein Halogenatom oder eine Hydrocarbonylsulfonyloxygruppe überführt wird.

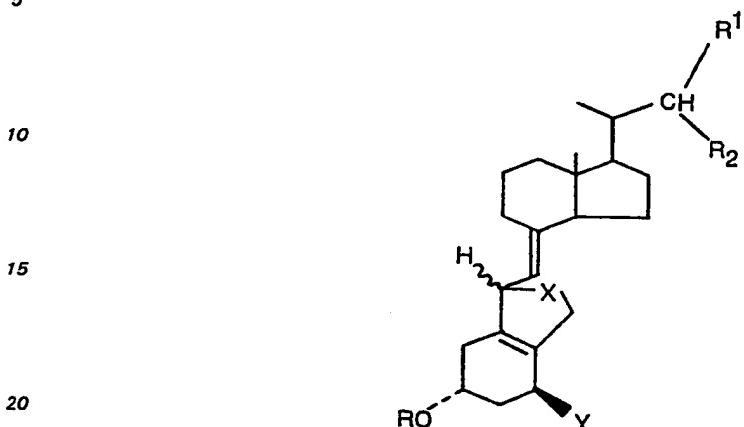
- 60 6. Verfahren gemäß einem der vorhergehenden Ansprüche, dadurch gekennzeichnet, daß ein Produkt erhalten wird, worin R¹ ein Halogenatom oder eine Hydroxyl- oder Hydrocarbonylsulfonyloxygruppe bedeutet, wodurch das Produkt in ein Produkt überführt wird, worin R¹ eine Gruppe der Form —ZR³, wie in Anspruch 1 definiert, anders als eine Hydroxylgruppe, darstellt.

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R revendications pour les Etats contractants: BE CH DE FR GB IT LI LU NL SE

1. Composés répondant à la formule générale

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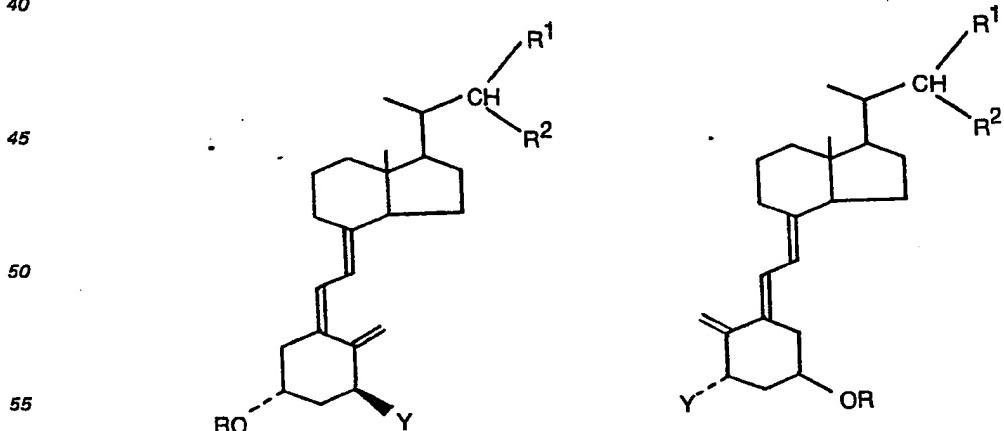


10 dans laquelle R représente un atome d'hydrogène ou un groupe protégeant la fonction hydroxyle, Y
 15 représente un atome d'hydrogène ou un radical hydroxyle éventuellement protégé, X représente un radical
 ---SO_2 ou le reste d'un diacylazodiénophile et R¹ représente un atome d'halogène, un radical hydrocarbylsulfonyloxy ou un groupe de la formule ---Z---R^3 (où Z représente ---O--- , ---S--- , ---SO--- , ---NR^4 ou
 20 $\text{---CR}^4\text{R}^5$ et R³, R⁴ et R⁵, qui peuvent être identiques ou différents, représentent chacun un atome
 25 d'hydrogène ou un radical aliphatique à chaîne droite ou à chaîne ramifiée, possédant de 1 à 12 atomes de
 30 carbone et qui peut éventuellement porter un ou plusieurs substituants) et R² représente un atome
 35 d'hydrogène, ou bien R¹ et R² représentent ensemble un radical oxo ou un groupe alkylidène
 éventuellement substitué, à l'exception que R¹ et R² ne forment pas, ensemble avec le groupe
 $\text{---CH(CH}_3\text{)CH---}$ auquel ils sont attachés, un radical possédant le squelette de la chaîne latérale 17 β -hydrocarbylique ramifiée de la vitamine D₂ ou de la vitamine D₃.

35 2. Composés suivant la revendication 1, caractérisés en ce que le diénophile est un composé
 diacylazoïque cyclique.

3. Composés des formules générales IV et IVa

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IVa (cis)

IV (trans)

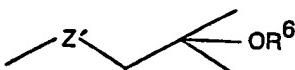
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65 dans lesquelles R, Y, R¹ et R² possèdent les significations qui leur ont été précédemment attribuées dans la revendication 1.

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4. Composés des formules générales I, IV et IVa, suivant l'un quelconque des revendications 1 à 3, dans lesquelles R¹ représente un atome d'halogène, un radical hydroxyle ou tosyloxy, ou un groupe de la formule

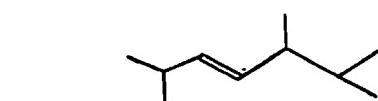
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10 (dans laquelle Z' représente —O—, —S—, —NH— ou —SO— et R⁶ représente un atome d'hydrogène ou un radical protégeant la fonction hydroxyle) et R² représente un atome d'hydrogène, ou bien R¹ et R² représentent ensemble un groupe alkylidène possédant de 1 à 8 atomes de carbone, éventuellement substitué par un ou plusieurs substituants choisis parmi les atomes d'halogènes et les radicaux hydroxyle éventuellement protégés.

15 5. Procédé de préparation de composés de la formule générale I suivant la revendication 1, dans laquelle R¹ et R² représentent ensemble un groupe oxo, caractérisé en ce que l'on soumet un composé de la formule III

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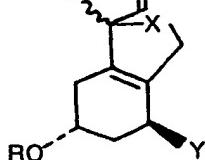


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(III)

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(dans laquelle R, Y et X possèdent les significations qui leur ont été précédemment attribuées dans la revendication 1) à une scission oxydante.

40 6. Procédé suivant la revendication 5, caractérisé en ce que l'on réduit ensuite l'aldéhyde de la formule I ainsi formé de façon à obtenir un composé de la formule I dans laquelle R¹ représente le radical hydroxyle.

7. Procédé suivant la revendication 5, caractérisé en ce que l'on fait ensuite réagir l'aldéhyde de la formule I ainsi formé sur un réactif de Wittig de manière à obtenir un composé de la formule I dans laquelle R¹ et R² représentent ensemble un groupe alkylidène éventuellement substitué dont la double liaison peut ensuite être réduite si on le souhaite.

45 8. Procédé de préparation de composés de la formule IV ou de la formule IVa telles que définies dans la revendication 3, caractérisé en ce que l'on déprotège un composé correspondant de la formule I telle que définie dans la revendication 1 par l'enlèvement du résidu X et ensuite, éventuellement après conversion du groupe R¹ en un autre groupe R¹, on isomérisé le composé de la formule IV ainsi obtenu en un composé de la formule IVa.

50 9. Procédé suivant la revendication 6 ou la revendication 8, caractérisé en ce que l'on obtient un produit dans lequel R¹ représente un radical hydroxyle et on convertit le radical hydroxyle en question en un atome d'halogène ou un radical hydrocarbysulfonyloxy.

55 10. Procédé suivant l'une quelconque des revendications 6, 8 et 9, caractérisé en ce que l'on obtient un produit dans lequel R¹ représente un atome d'halogène ou un radical hydroxyle ou hydrocarbysulfonyloxy, produit que l'on convertit ensuite en une substance dans laquelle R¹ représente un groupe de la formule —ZR³ telle que définie dans la revendication 1, autre qu'un radical hydroxyle.

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Revendications pour l'Etat contractant AT:

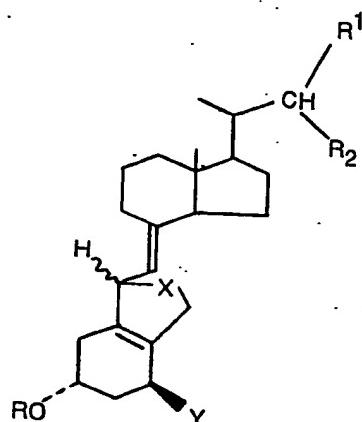
1. Procédé de préparation d'composés de la formule générale

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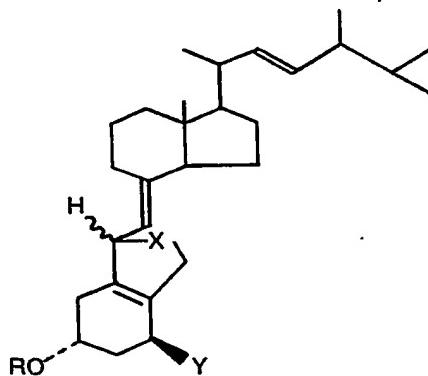


(dans laquelle R représente un atome d'hydrogène ou un groupe protégeant la fonction hydroxyle, Y
 25 représente un atome d'hydrogène ou un radical hydroxyle éventuellement protégé, X représente un radical —SO₂ ou le reste d'un diacylazodiénophile et R¹ représente un atome d'halogène, un radical hydrocarbysulfonyloxy ou un groupe de la formule —Z—R³ (où Z représente —O—, —SO—, —SO—, —NR⁴— ou —CR⁴R⁵— et R³, R⁴ et R⁵, qui peuvent être identiques ou différents, représentent chacun un atome d'hydrogène ou un radical aliphatique à chaîne droite ou à chaîne ramifiée, possédant de 1 à 12 30 atomes de carbone et qui peut éventuellement porter un ou plusieurs substituants) et R² représente un atome d'hydrogène, ou bien R¹ et R² représentent ensemble un radical oxo ou un groupe alkylidène éventuellement substitué, à l'exception que R¹ et R² ne forment pas, ensemble avec le groupe —CH(CH₃)CH— auquel ils sont attachés, un radical possédant le squelette de la chaîne latérale 17β-hydrocarbylique ramifiée de la vitamine D₂ ou de la vitamine D₃) caractérisé en ce que l'on soumet un 35 composé de la formule III

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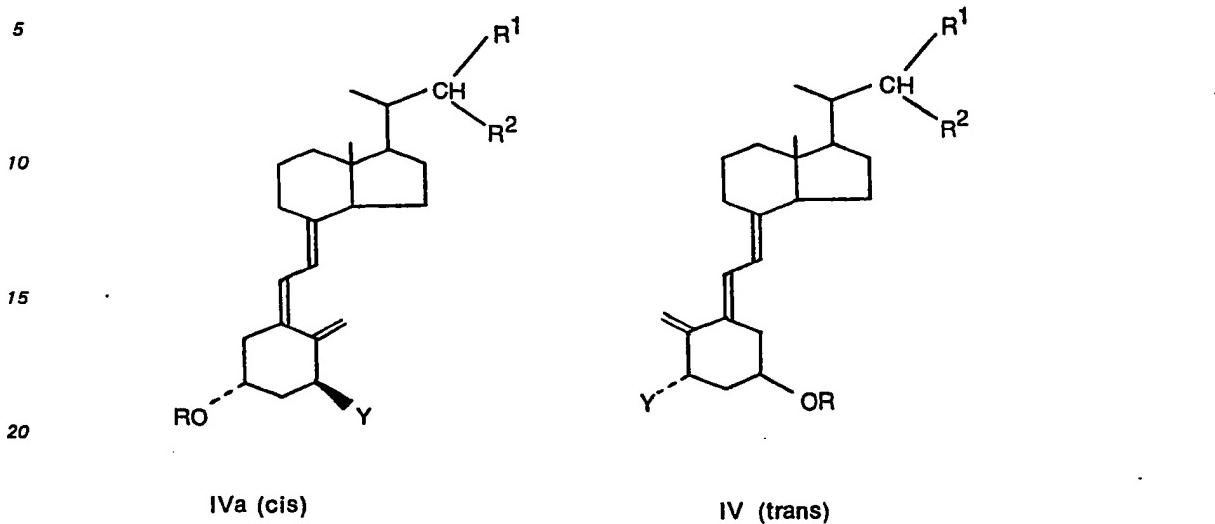
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(dans laquelle R, Y et X possèdent les significations qui leur ont été précédemment attribuées) à une 55 scission oxydante de manière à former un aldéhyde de la formule I (dans laquelle R¹ et R² représentent ensemble un radical oxo) et ensuite, si on le souhaite, on fait réagir l'aldéhyde en question de la formule I sur un réactif de Wittig de manière à obtenir un composé de la formule I dans laquelle R¹ et R² représentent ensemble un radical alkylidène éventuellement substitué dont on peut ensuite réduire la double liaison si on le souhaite, ou bien on réduit l'aldéhyde en 60 question de la formule I de façon à obtenir un composé de la formule I dans laquelle R¹ représente un radical hydroxyl, le radical hydroxyle étant ensuite éventuellement converti en un atome d'halogène, un radical hydrocarbysulfonyloxy ou un groupe de la formule —ZR³ t il que précédemment définie autre qu'un radical hydr oxyde.

2. Procédé suivant la revendication 1, caractérisé en ce que le diénophile est un composé diacylazoïque 65 cyclique.

3. Procédé de préparation de composés de la formule général IV ou de la formule générale IVa



- (dans lesquelles R, Y, R¹ et R² possèdent les significations qui leur ont été précédemment attribuées dans la revendication 1), caractérisé en ce que l'on déprotège un composé correspondant de la formule I telle que définie dans la revendication 1 par l'enlèvement du reste X et ensuite, éventuellement après conversion du groupe R¹ en un autre groupe R¹, on isomérise le composé de la formule IV ainsi obtenu en un composé de la formule IVa.
4. Procédé suivant l'une quelconque des revendications précédentes de préparation de composés des formules générales I, IV ou IVa dans lesquelles R¹ représente un atome d'halogène, un radical hydroxyle ou tosyloxy, ou un groupe de la formule

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- 40 (dans laquelle Z' représente —O—, —S—, —NH— ou —SO— et R⁶ représente un atome d'hydrogène ou un radical protégeant la fonction hydroxyle) et R² représente un atome d'hydrogène, ou bien R¹ et R² représentent ensemble un radical alkylidène possédant de 1 à 8 atomes de carbone, éventuellement substitué par un ou plusieurs substituants choisis parmi les atomes d'halogène et les radicaux hydroxyle éventuellement protégés.
- 45 5. Procédé suivant l'une quelconque des revendications précédentes, caractérisé en ce que l'on obtient un produit dans lequel R¹ représente un radical hydroxyle et on convertit le radical hydroxyle en question en un atome d'halogène ou un radical hydrocarbysulfonyloxy.
- 50 6. Procédé suivant l'une quelconque des revendications précédentes, caractérisé en ce que l'on obtient un produit dans lequel R¹ représente un atome d'halogène ou un radical hydroxyle ou hydrocarbysulfonyloxy, produit que l'on convertit ensuite en un substance dans laquelle R¹ représente un groupe de la formule —ZR³ telle que définie dans la revendication 1 autre qu'un radical hydroxyle.

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